

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
6 June 2002 (06.06.2002)

PCT

(10) International Publication Number  
**WO 02/43801 A2**

(51) International Patent Classification<sup>7</sup>: **A61N**  
(21) International Application Number: PCT/US01/43237  
(22) International Filing Date:  
19 November 2001 (19.11.2001)  
(25) Filing Language: English  
(26) Publication Language: English  
(30) Priority Data:  
60/252,946 24 November 2000 (24.11.2000) US  
(71) Applicant (for all designated States except US): **U-SYS-TEMS, INC.** [US/US]; 110 Rose Orchard Way, San Jose, CA 95134 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

**Published:**

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(72) Inventor; and  
(75) Inventor/Applicant (for US only): **WANG, Shih-Ping** [US]; 409 Becker Lane, Los Altos, CA 94022 (US).  
(74) Agent: **KAVRUKOV, Ivan, S.**; Cooper & Dunham LLP, 1185 Avenue of the Americas, New York, NY 10036 (US).

(54) Title: METHOD AND SYSTEM FOR USING BREAST ULTRASOUND INFORMATION TO FACILITATE BREAST CANCER SCREENING

(57) **Abstract:** A breast cancer screening system having a display apparatus that simultaneously displays a succession of ultrasound image slices, the ultrasound image slices being taken from successive planes in a breast volume substantially parallel to a plane of a predetermined x-ray mammogram view of the breast, is described. The ultrasound image slices are preferably displayed adjacent to a display of the x-ray mammogram view itself, such that a screening radiologist may elect to view the ultrasound image slices simultaneously with the x-ray mammogram view. Preferably, the predetermined x-ray mammogram view is a standardized x-ray mammogram view such as the craniocaudal view (CC) or mediolateral oblique view (MLO). The screening system may also perform computer-aided diagnosis (CAD) algorithms on the ultrasound image slices and/or the x-ray mammogram view. A breast ultrasound adapter is also described for facilitating reliable acquisition of the breast ultrasound scans. The breast ultrasound adapter optionally comprises acoustic transducers for vibrating the breast tissue such that vibrational resonance information is acquired in addition to the primary ultrasound information. The vibrational resonance information is provided to the CAD system in conjunction with the primary ultrasound information and the x-ray mammogram information. Lesion features extracted from the vibrational resonance information are fed to a classifier, *e.g.* an artificial neural network, in conjunction with features extracted from the primary ultrasound information and the x-ray mammogram information. The x-ray mammogram, the ultrasound image slices, the vibrational resonance images, and the CAD output can be simultaneously displayed in selected combinations according to input commands from the user.



WO 02/43801 A2

**METHOD AND SYSTEM FOR USING BREAST  
ULTRASOUND INFORMATION TO FACILITATE  
BREAST CANCER SCREENING**

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Shih-Ping Wang

**CROSS-REFERENCE TO RELATED APPLICATIONS**

This application claims the benefit of United States Provisional Application No. 60/252,946, filed November 24, 2000, which is incorporated by reference herein.

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**FIELD**

The present invention relates to medical imaging systems adapted to facilitate breast cancer screening. In particular, the present invention relates to the manipulation and display of breast ultrasound information in a manner that is convenient, amenable to standardization, and readily usable by radiologists accustomed to traditional x-ray mammogram analysis.

15

**BACKGROUND**

Breast cancer is the most common cancer among women other than skin cancer, and is the second leading cause of cancer death in women after lung cancer. The American Cancer Society currently estimates that there are about 184,200 new cases of invasive breast cancer per year among women in the United States and 41,200 deaths per year from the disease. Prevention and early diagnosis of breast cancer are of foremost importance. Because early breast cancer does not produce symptoms, the American Cancer Society recommends a mammogram and a clinical breast examination every year for women over the age of 40.

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X-ray mammography is currently the imaging method for mass screening of breast cancer. In health maintenance organizations (HMO's) and other medical organizations, specialized x-ray mammography clinics designed for high patient throughput are being increasingly used to screen as many women as possible in a time and cost efficient manner. X-ray mammography practice in the United States has become largely standardized. For each x-ray mammogram screening of a patient, two standard x-ray mammogram views of each breast are commonly taken: a top (head-to-toe) view ordinarily called the

craniocaudal view ("CC"), and a lateral view ordinarily called the mediolateral oblique view ("MLO"). Several efficiencies arise by virtue of this standardization, including: the ability for examinations to be conducted by x-ray technicians; the ability of a radiologist to analyze mammograms without being physically present as the x-rays are taken; the ability  
5 to compare and statistically track large numbers of x-ray mammograms taken from different facilities; the ability to track changes in a single patient over time even if the x-ray mammograms are taken at different facilities; the ability of radiologists to gain recursive expertise in analyzing the standard x-ray mammogram views; and the repeatability of results.

10 The standardization of x-ray mammograms also yields benefits in the public health care area, including the ability for the U.S. government to provide a fixed and predictable per-mammogram reimbursement for Medicare patients. Additionally, HMO's and other medical insurers are provided with predictable outlays for breast cancer screening of their member patients using x-ray mammography.

15 A well-known shortcoming of x-ray mammography practice, however, is found in the case of dense-breasted women including patients with high content of fibroglandular tissues in their breasts. Because fibroglandular tissues have higher x-ray absorption than the surrounding fatty tissues, portions of breasts with high fibroglandular tissue content are not well penetrated by x-rays and thus the resulting mammograms would contain little or  
20 no information in areas where fibroglandular tissues reside. A recent study by Lehman et. al. on 46,340 patients, published in the December 1999 issue of the American Journal of Roentgenology (AJR), reports that the proportion of dense breasts (summing those with "heterogeneously dense" and "extremely dense" breasts) account for about 52% of women with age range of 35-39, 47% of age range 40-49, 32% of age range 50-59, 24% of age  
25 range 60-69, 23% of age range 70 or older, and 36% for all ages. Lehman et. al., "Effect of Age and Breast Density on Screening Mammograms with False-Positive Findings," AJR 173:1651-1655 (December 1999). For the estimated 36% of the female population who have dense breasts, this means that at least a portion of the breast area on the x-ray mammogram cannot be scrutinized for lesions by x-ray mammography alone. As a result,  
30 lesions camouflaged by dense breast tissue may go undetected. Indeed, a study by Kolb et al on 18,005 consecutive patients, as reported in Jalali, "Sound Combination: Ultrasound

Paired With a Mammography Can Improve Cancer Detection for Dense-Breasted Women,” ADVANCE for Administrators In Radiology and Radiation Oncology, pp. 68-70, states that x-ray mammography were able to detect only 56 cancers of the 80 cancers in 7,202 patients with dense breasts.

- 5           It has been found that the use of ultrasound mammography (sonomammography) in conjunction with conventional x-ray mammography can drastically increase the early breast cancer detection rate. Ultrasound mammograms display the acoustic impedance of individual slices of breast tissue, and therefore may allow detection of breast lesions where x-ray mammography alone fails. The study by Kolb, *supra*, revealed this drastic increase.
- 10   In a study of 18,005 consecutive screening examinations on 10,200 women who had mammograms, ultrasound exams, and physical examination during a three-year period at Dr. Kolb’s New York City office, 7,202 had dense breasts. Among that group, 80 breast cancers were detected in 75 women. Mammography alone detected 56 of the cancers (70 percent), while ultrasound in addition to mammography detected 75 of the cancers (94
- 15   percent). The remaining five cancers (6 percent) were found through physical exam. All cancers were confirmed with surgical biopsy. Thus, the use of ultrasound mammography increased the early breast cancer detection rate by 33 percent as compared with x-ray mammography alone.

- Vibrational resonance techniques, such as those discussed in U.S. Pats. 5,919,139
- 20   and 6,068,597, have been proposed for analyzing suspect tumors. As discussed in Lowers, J., “Experimental Modes Abound For Detecting Breast Cancer: Vibrational Resonance Technique Among the Contenders,” Women’s Health Supplement to Diagnostic Imaging (April 2001) at pp. 15-17, an audio speaker is attached to the ultrasound probe to introduce audio-range vibrational tones (*e.g.*, 69-247 Hz) into the patient during the acquisition of
- 25   power Doppler ultrasound frames. Different tissue types often vibrate by different amounts responsive to the acoustic signals, and the different vibrations result in different power Doppler readings. It has been found that the vibrations can help to clearly differentiate suspect lesions, even those that appear isoechoic with surrounding tissue (*i.e.*, are less noticeable) on B-scan ultrasound frames. However, the Lowers reference *supra* is
- 30   directed to the analysis of known suspicious lesions rather than to initial breast cancer screening environments.

It has been found that a major inhibitor to the use of ultrasound mammography in breast cancer screening is the difficulty of integrating it into today's specialized, high-throughput x-ray mammography environment. This is especially important in light of the substantial training and expertise built up over several years in physicians reading the x-ray  
5 mammograms and in x-ray technicians conducting standardized x-ray mammography examinations.

An additional major inhibitor to the use of ultrasound in breast cancer screening is the additional physician time that would be required to conduct the ultrasound examination in addition to analyze the ultrasound. In the current health-care environment, physician  
10 time is a crucial concern. Additional physician diagnosis time per patient means additional cost. Efficacy of radiological procedures is today measured by the cost in dollars per quality adjusted life year (QALY), with procedures costing more than \$100,000 per QALY being neither encouraged nor prescribed. It would be desirable to provide a screening system that integrates ultrasound mammography into current breast cancer screening  
15 methodologies without requiring technological upheaval, without requiring substantial additional physician time to conduct the examination and to interpret the ultrasound data, and without losing the benefits of the many years of professional expertise developed in relation to current x-ray mammography methods. Thus, it has been found that perhaps the most valuable and saleable feature of a screening system is ready and efficient usability by  
20 today's medical professionals.

Various methods have been proposed for using ultrasound information in breast cancer screening. U.S. Pat. 5,938,613 to Shmulewitz ("the '613 patent"), which is incorporated by reference herein, discusses a method and apparatus for performing sonomammography and enhanced x-ray imaging in which ultrasound equipment is  
25 integrated with mammography equipment to generate ultrasonic images of the internal structure of breast tissue that are in geometric registration with an x-ray mammogram. However, the '613 patent is primarily directed to solving registration problems between the ultrasound and x-ray mammograms, and does not disclose methods for viewing and analyzing the data that are sufficiently suitable for everyday practice by radiologists, and in  
30 particular for radiologists that are accustomed to viewing standard CC and MLO views of x-ray mammograms in their everyday practice. Additionally, significant cost and

complexity are devoted to the registration of the ultrasound data and the x-ray mammography data, whereas it has been found that useful analysis can be performed without requiring such precise registration.

U.S. Pat. 5,984,870 to Giger et al. ("the '870 patent"), which is incorporated by  
5 reference herein, discusses a method and apparatus for the computerized automatic analysis of lesions in ultrasound images. Geometric features associated with cancerous conditions may be extracted from an ultrasound mammogram. Features of the lesion from the ultrasound mammogram can then be fed into an artificial neural network together with features of the lesion from an x-ray mammogram and/or an MRI to derive an estimate of  
10 the likelihood of malignancy. However, the '870 patent does not disclose a manner of presenting the ultrasound data itself to the radiologist such that effective results checking and ultimate diagnosis can occur. It is believed that a key question relevant to the effectiveness and saleability of an ultrasound mammography system, a question that appears to be regularly avoided by the prior art, is whether today's doctors will want to use  
15 the system in everyday screening, and whether today's breast cancer screening clinics will want to buy the system.

It would be desirable to provide a breast cancer screening system that increases early breast cancer detection by readily incorporating ultrasound mammography into today's existing breast cancer screening infrastructure.

20 It would be still further desirable to provide a screening system in which ultrasound mammography results can be integrated with x-ray mammography results without losing the benefits of the many years of professional expertise developed in relation to current standardized x-ray mammography methods.

It would be still further desirable to provide a screening system that takes little or  
25 no special familiarization or training from the radiologist in order to effectively view ultrasound information in combination with the x-ray mammogram information.

It would be still further desirable to provide a screening system that provides standardized, repeatable, and easily interpretable ultrasound mammography results that may be quickly analyzed in combination with x-ray mammogram information such that  
30 screening cost per patient is not substantially increased.

It would be still further desirable to provide a screening system wherein upon acquisition of the system, any increase in breast cancer screening costs is offset by savings brought about by an increased early breast cancer detection rate, whereby cost per patient QALY is ultimately reduced.

- 5           It would be even further desirable to provide an improved method of using combinations of x-ray mammography-based and ultrasound-based data in a computer-aided diagnosis (CAD) system.

- It would be still further desirable to provide a screening system that provides standardized, repeatable, and easily interpretable ultrasound mammography results and  
10   ultrasound computer-aided diagnosis (CAD) results that may be quickly analyzed in combination with x-ray mammogram information such that screening cost per patient is not substantially increased.

#### SUMMARY

- 15           In accordance with a preferred embodiment, a screening system is provided having a display apparatus that simultaneously displays a succession of ultrasound image slices, the ultrasound image slices being taken from successive planes in a breast volume substantially parallel to a plane of a predetermined x-ray mammogram view of the breast. The ultrasound image slices are preferably displayed adjacent to a display of the x-ray  
20   mammogram view itself, such that a screening radiologist may view the ultrasound image slices simultaneously with the x-ray mammogram view. The display of ultrasound image slices may be used to inspect areas of dense breast tissue that appear opaque on the x-ray mammogram view. Because they are taken from planes in the breast parallel to the x-ray mammogram view plane, the ultrasound image slices are of immediate and familiar  
25   significance to the screening radiologist. The breast cancer screening process is thereby made more thorough while at the same time being about as time-efficient as a screening of the x-ray mammogram view alone. Preferably, the predetermined x-ray mammogram view is a standardized x-ray mammogram view such as the craniocaudal view (CC) or mediolateral oblique view (MLO), whereby further advantages of standardization may be  
30   realized, including the ability to share, store, and track the ultrasound image slices in a manner similar to the way x-ray mammogram images are shared, stored, and tracked.

According to a preferred embodiment, ultrasound scans are taken of the breast when flattened or compressed along a plane substantially parallel to the x-ray mammogram view plane. The ultrasound scans are taken by an ultrasound probe, which in a preferred embodiment scans the flattened breast in planes perpendicular to the x-ray mammogram view plane. In such case the screening system is adapted to reconstruct a 3-dimensional volume of the acoustic impedance of the breast, and then interpolate within the 3-dimensional volume to compute the ultrasound image slices along planes substantially parallel to the x-ray mammogram view plane. It is within the scope of the preferred embodiments, however, for the ultrasound probe to scan the flattened breast in planes parallel to the x-ray mammogram view plane, wherein the ultrasound image slices are directly computed from the ultrasound scan data without the need for 3-dimensional volume reconstruction.

According to another preferred embodiment, a breast ultrasound adapter is provided for facilitating the reliable acquisition of breast ultrasound scans. The breast ultrasound adapter comprises an open fluid reservoir defined by side walls and a bottom membrane for contacting a breast surface. The breast ultrasound adapter is designed for placement on the breast of a supine patient such that it can be filled with water and/or a water-containing polymeric gel or other suitable fluid until the bottom membrane is submerged. The bottom of an ultrasound probe is submerged in the fluid, and the probe is swept in a lateral direction across the breast, the bottom of the probe remaining submerged during the lateral sweep. Preferably, the bottom of probe is maintained in a plane parallel to the fluid surface and does not contact the bottom membrane during the lateral sweep. Advantageously, the breast is maintained in a substantially consistent position and in a consistent state of compression during the scanning process, thereby facilitating consistency among the ultrasound slices forming the three-dimensional image volume. Water, water-containing polymeric gel, and/or oil are preferably used to improve the contact between the bottom membrane and the breast. Optionally, acoustic transducers such as audio speakers are fixed to the breast ultrasound adapter for use in vibrational resonance applications. In one preferred embodiment, the ultrasound probe is swept by hand across the top surface while position sensors detect its position. Alternatively, a



mechanically driven probe is coupled to the top of the breast ultrasound adapter, the probe position being sensed by mechanical transducers.

According to another preferred embodiment, the screening system performs computer-aided diagnosis (CAD) algorithms on the ultrasound image slices for detecting  
5 suspicious locations in the breast. Highlighting marks are superimposed at the suspicious locations of the ultrasound image slices, or alternatively an additional display device is used to display suspicious lesion information using other methods, *e.g.*, using a three-dimensional display method. Optionally, the screening system includes devices that perform CAD algorithms on the x-ray mammogram view as well. The x-ray mammogram,  
10 the ultrasound image slices, the ultrasound CAD outputs, and the x-ray mammogram CAD outputs may be quickly displayed to the user in various combinations according to input commands from the user.

According to another preferred embodiment, additional ultrasound information is acquired using vibrational resonance techniques. For each primary ultrasound slice being  
15 displayed to the user, one or more additional corresponding vibrational resonance images is superimposed thereon or provided in an additional display. Optionally, the user may activate a toggle switch to cyclically superimpose/remove the vibrational resonance image on/from the primary ultrasound slice. In another preferred embodiment, the vibrational resonance information is provided to the CAD system in conjunction with the x-ray  
20 mammogram and/or the primary ultrasound scan data. Vibrational resonance-based features of a lesion are extracted from the vibrational resonance image, and then fed to a classifier, *e.g.* an artificial neural network, in conjunction with features extracted from the x-ray mammogram and/or primary ultrasound images, for determining a likelihood of malignancy or other metric of suspiciousness.

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### BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 illustrates steps taken in accordance with a preferred embodiment;

FIG. 2 illustrates a diagram of an ultrasound probe scanning a breast that is flattened along an x-ray mammogram view plane in accordance with a preferred  
30 embodiment;

FIG. 3 illustrates a conceptual diagram of 3-dimensional breast volume reconstruction and ultrasound image slice interpolation in accordance with a preferred embodiment;

FIG. 4 illustrates a user display of a breast cancer screening system in accordance  
5 with a preferred embodiment;

FIG. 5 illustrates an ultrasound image slice display of the user display of FIG. 4;

FIG. 6 illustrates steps taken in accordance with a preferred embodiment;

FIG. 7 illustrates a user display of a breast cancer screening apparatus in  
accordance with a preferred embodiment;

10 FIG. 8 illustrates steps taken in accordance with a preferred embodiment;

FIG. 9 illustrates a user display of a breast cancer screening apparatus in  
accordance with a preferred embodiment;

FIG. 10 illustrates steps taken in accordance with a preferred embodiment;

FIG. 11 illustrates steps taken in accordance with a preferred embodiment;

15 FIGS. 12A and 12B illustrate steps taken in accordance with a preferred  
embodiment;

FIG. 13 illustrates a breast ultrasound adapter for facilitating breast ultrasound  
scans in accordance with a preferred embodiment; and

FIG. 14 illustrates a partially automated apparatus for performing breast ultrasound  
20 scans in accordance with a preferred embodiment.

## DETAILED DESCRIPTION

FIG. 1 illustrates steps taken in accordance with a preferred embodiment in which  
ultrasound data is used in conjunction with x-ray mammogram data for a more thorough  
25 diagnosis as compared to x-ray mammogram data alone. At step 102, an x-ray  
mammogram view plane is chosen. More than one x-ray mammogram view plane may be  
chosen, it being understood that the steps of FIG. 1 are to be repeated for each chosen x-  
ray mammogram view plane. As used herein, an x-ray mammogram view plane is a plane  
in which x-ray film is placed for receiving x-ray radiation in an x-ray mammogram view.  
30 The x-ray mammogram view plane may be described using any numerical criterion  
sufficient to describe a plane, for example, the x-y-z coordinates of the tip of a unit vector

normal to the plane. For well-known standardized x-ray mammogram views, such as the CC and MLO views, the x-ray mammogram view plane can be identified by a simple reference to that x-ray mammogram view, it being understood that persons skilled in the art would recognize the proper position and orientation of the plane.

- 5           At step 104, an x-ray mammogram is obtained for the chosen x-ray mammogram view plane. Generally speaking, step 104 is performed using standard x-ray mammography methods at a breast cancer screening clinic. Alternatively, in the case where the steps of FIG. 1 are being carried out several hours, days, or weeks after all the clinical measurements are taken, step 104 is performed by accessing the appropriate
- 10       database or accessing the appropriate physical x-ray film containing the x-ray mammogram for the chosen x-ray mammogram view plane. At step 104, the x-ray mammogram is typically a x-ray film acquired with a film-screen system. However, the x-ray mammogram may also be acquired by a digital detector, such as that used in a digital mammography system, in the form of a digital mammogram.
- 15           At step 106, the breast is flattened along an ultrasound-absorbing plate placed in the chosen x-ray mammogram view plane, and at step 108 ultrasound scan data on the flattened breast is obtained. At step 110, a three-dimensional volume reconstruction is performed using the ultrasound scan data to form a three-dimensional acoustic impedance map of the breast. At step 112, ultrasound image slices are interpolated along planes
- 20       parallel to the chosen x-ray mammogram view plane using the three-dimensional acoustic impedance map. Finally, at step 114 the x-ray mammogram and the ultrasound image slices for the chosen x-ray mammogram view plane are simultaneously displayed for viewing by the screening radiologist.

- Importantly, it is to be appreciated that step 106, together with subsequent step
- 25       108, is not required to be performed at the same time, on similarly calibrated equipment, or even at the same screening facility as step 104. Indeed, in many instances it may be more convenient to perform examinations on separate machines or in separate facilities to increase patient throughput. Advantageously, in accordance with the preferred embodiments, precise registration between the ultrasound image slices and the
- 30       corresponding x-ray mammogram views is not required. It is only preferred that the breast be flattened against a plane substantially parallel to the x-ray mammogram view plane, so

that when ultrasound image slices are derived from the ultrasound scan data, they will generally correspond to the chosen x-ray mammogram view.

FIG. 2 illustrates a diagram corresponding to steps 106 and 108 of FIG. 1, wherein an ultrasound probe 202 scans a breast 204 that is flattened along an ultrasound-absorbing plate 206. The ultrasound-absorbing plate 206 lies in the chosen x-ray mammogram view plane, which for the example of FIG. 2 is the x-y plane in the axis system shown. Ultrasound scan data is obtained using methods known in the art. The breast 204 may be flattened either by hand, or may be flattened by a planar device that is integral with an ultrasound probe, such as that described in U.S. Pat. 5,938,613, *supra*. In the case of a planar ultrasound probe, an impedance-matching ultrasound gel or gel bag should be placed between the breast and the probe for best results. Because typical hand-held ultrasound probes are about 38 mm wide, it may be desirable to incorporate two or more parallel probes to cover the breast in one scan. Or alternatively, an automatic raster scan mechanism may be used to cover the breast. The dynamic range of the ultrasound data samples is chosen such that approximately 8 bits of gray scale resolution is yielded in the ultrasound image slices.

Also shown in FIG. 2 are fixed sensors 208 and 210, which together with an electromagnetic transmitter embedded into ultrasound probe 202 form an electromagnetic position sensing system. The electromagnetic position sensing system provides the necessary mathematical link between the acoustic reflection data obtained by the ultrasound probe 202 and the corresponding position within the breast 204. Suitable electromagnetic sensing systems that may be used in accordance with the preferred embodiments are described in Cheng et al, "Automated Detection of Breast Tumors in Ultrasonic Images Using Fuzzy Reasoning," *Proceedings of the IEEE Computer Society International Conference on Image Processing Volume III*, pp. 420-423 (October 26-29, 1997) (hereinafter "Cheng"), the contents of which are hereby incorporated by reference. Other types of ultrasound probe position sensing systems are known in the art, and any of a variety of such systems may be used in accordance with the preferred embodiments, including servo-mechanical sensing systems (*see, e.g.*, U.S. Pat. 5,433,202 to Mitchell et. al., the contents of which are hereby incorporated by reference) and fixed plate scanning systems (*see, e.g.*, U.S. Pat. 5,640,956 to Getzinger et. al., the contents of which are hereby

incorporated by reference). Alternatively, inertial-type of positional sensors, such as those used in the Hewlett Packard CapShare 920 Portable E-Copier hand-held page scanner, may also be used.

Preferably, the ultrasound scans taken by ultrasound probe 202 are taken in planes  
5 that are normal to the chosen x-ray mammogram view plane (the x-y plane in FIG. 2). Thus, in FIG. 2 the ultrasound scans are taken for a series of planes parallel to the x-z plane, which is normal to x-y plane, resulting in a succession of two-dimensional ultrasound planes denoted herein as  $n_i$ , where "i" is a counter variable for the successive ultrasound planes. It is preferable to take the ultrasound scans along planes normal to the  
10 x-ray mammogram view plane because the resulting scans are normal to the flattened breast and therefore are of a shallow depth compared to ultrasound scans taken parallel to the x-ray mammogram view plane. Because of the shallower depth, greater ultrasound image resolution can be obtained. As a typical numerical example, ultrasound scans taken normal to a flattened breast would only need to penetrate about 4 cm of breast tissue,  
15 whereas ultrasound scans taken parallel to the plane of the flattened breast would need to penetrate about 15 cm of breast tissue. Due to acoustic attenuation in the human body that increases with frequency (a factor of about 0.6 dB/MHz-cm), lower frequency probes operating near 3 MHz would be required for taking ultrasound scans parallel to the plane of the flattened breast, with the use of lower probe frequency yielding a lower spatial  
20 resolution of about 0.1 cm. In contrast, higher frequency probes operating at about 7.5 MHz can be used for taking ultrasound scans normal to the flattened breast, with the use of the higher probe frequency yielding a better spatial resolution of about 0.05 cm.

However, although it is preferable ultrasound probe 202 scan in a direction normal to the chosen x-ray mammogram view plane, it is nevertheless within the scope of the  
25 preferred embodiments for scanning to occur parallel to the x-ray mammogram view plane in certain circumstances. This is because even though the resolution is diminished, one advantage of scanning parallel to the x-ray mammogram view plane is that less processing time is required to generate the ultrasound image slices, which are generally directly yielded from the ultrasound scans themselves. When the ultrasound scans are taken by the  
30 ultrasound probe 202 parallel to the x-ray mammogram view plane, the ultrasound image slices may be directly computed without the need for performing the three-dimensional

reconstruction and interpolation at step 112. Because the lower end of the target size lesions is usually about 0.5 cm, the lesser spatial resolution of 0.1 cm associated with the lower frequency probe may be sufficient in some clinical environments.

FIG. 3 illustrates a diagram corresponding to steps 110 and 112 of FIG. 1, in particular a conceptual diagram of 3-dimensional breast volume reconstruction and ultrasound image slice interpolation in accordance with a preferred embodiment. FIG. 3 conceptually shows a 3-dimensional acoustic impedance matrix 302 constructed from the plurality of two-dimensional ultrasound planes  $n_i$  resulting from step 108. The 3-dimensional acoustic impedance matrix 302 may be constructed from the ultrasound planes  $n_i$  using any of a variety of known 3-dimensional volume formation algorithms. FIG. 3 also conceptually shows a representative ultrasound image slice 304 resulting from the application of an interpolation algorithm to the 3-dimensional acoustic impedance matrix 302, the ultrasound image slice 304 being substantially parallel to the x-ray mammogram view plane (*e.g.*, the x-y axis in FIGS. 2 and 3), the ultrasound image slices being denoted herein by  $m_j$ , where  $j$  is a counter variable for the successive ultrasound image slices. The successive ultrasound image slices  $m_j$  may be computed using any of a variety of known 3-to-2 dimensional projection algorithms. Depending on the interpolation/projection algorithm used, each ultrasound image slice may represent instantaneous values along a single plane within the 3-dimensional acoustic impedance matrix. Alternatively, each ultrasound image slice may be a "thick" one representing an integration of the 3-dimensional acoustic impedance matrix over a predetermined thickness (*e.g.*, 0.5 cm). The latter algorithm is usually preferable because more data within the 3-dimensional acoustic impedance matrix is used to compute the actual result, and less data is "discarded" by the projection algorithm.

In accordance with a preferred embodiment, the successive ultrasound image slices  $m_j$  correspond to planes in the breast volume that are no greater than a predetermined spacing distance apart. The predetermined spacing distance is selected to reduce the probability of missed lesions due to undersampling along an axis perpendicular to the x-ray mammogram view plane (*e.g.*, the x-axis in FIGS. 2 and 3). At the same time, however, the predetermined spacing distance should not be so small as to cause the screening radiologist to view an untenably large number of ultrasound image slices. For one

preferred embodiment, a predetermined spacing distance of approximately 0.5 cm is suitable for many applications. However, it is also anticipated that some physicians may prefer higher spatial resolution (more slices with spacing less than 0.5 cm) and some may prefer lower spatial resolution (fewer slices with spacing more than 0.5 cm). In these cases,  
5 more or fewer slices will be provided to suit the physician's taste.

FIG. 4 illustrates a user display 400 of a breast cancer screening system in accordance with a preferred embodiment. User display 400 comprises an x-ray mammogram display 402 for showing a chosen x-ray mammogram view 404. The x-ray mammogram display 402 may comprise a TV monitor for displaying a digitized form of  
10 the x-ray mammogram view, or alternatively may comprise a mechanical display such as a conveyor-based system and light box for displaying x-ray mammogram film. User display 400 further comprises an ultrasound slice display 406 for simultaneously displaying a film containing a plurality of ultrasound image slices  $m_j$  adjacent to one another, the display 406 being positioned such that it may be viewed simultaneously with the x-ray  
15 mammogram display 402. The ultrasound slice display 406 of FIG. 4 shows a display of eight ultrasound image slices  $m_j$ , although as few as four slices and as many as 16 or more slices may be displayed depending on screening radiologist desires. When displaying a digital mammogram on a TV monitor, it is sometimes preferred to also display the ultrasound slices on the same or an adjacent TV monitor. Or, while the x-ray mammogram  
20 may be in the film form, the ultrasound slices be displayed on an adjacent TV monitor. Alternatively, while the x-ray mammogram may be displayed on a TV monitor, the ultrasound slices be displayed on a film on an adjacent light box.

User display 400 may further comprise an optional keyboard 408, an optional mouse 410, and a plurality of optional control knobs/switches 412 for controlling the  
25 brightness, contrast, zoom, and other functions that may be installed with the user display 400. An example of an x-ray mammogram display apparatus that may be used in conjunction with the preferred embodiments can be found in U.S. Pat. 5,917,929 to Marshall et. al., the contents of which are hereby incorporated by reference.

The user display 400, and in particular the ultrasound slice display 406, may be  
30 advantageously used by the screening radiologist to quickly determine if there are problematic areas of dense breast tissue that would otherwise be camouflaged in the x-ray

mammogram view 404. A conceptual diagram of this is shown in FIG. 4, in which a spherically-shaped lesion 414 does not show up on the x-ray mammogram view 404 because it is camouflaged by an area of dense fibroglandular breast tissue 416. The lesion 414 is shown as a dotted line in x-ray mammogram view 404 to signify that the radiologist cannot see it. In accordance with a preferred embodiment, however, the lesion 414 does indeed show up among the plurality of ultrasound image slices  $m_j$  in ultrasound slice display 406, appearing in three of the ultrasound image slices as areas marked 414a, 414b, and 414c in FIG. 4. Thus, the use of the ultrasound slice display 406 in ready conjunction with the x-ray mammogram view 404 can avoid missed diagnoses due to camouflage of lesions within dense breast tissue.

Conversely, another advantage in accordance with the preferred embodiments can avoid increased false positive screening results due to dense breasts. In particular, (*see* Lehman et. al., *supra*), dense breasts typically lower the specificity (increased false positives) of x-ray screening examinations. A lesion may be obscured by the dense fibroglandular tissue or an apparent lesion on the x-ray mammogram view may in fact be due to a coincidental alignment of unrelated breast tissue that appears as a single lesion in the x-ray mammogram view. Such occurrences would readily be revealed by a quick examination of the ultrasound image slices, wherein the screening radiologist would quickly determine that there is no single lesion in the ultrasound image slices but rather a plurality of unrelated tissue segments. Especially in the circumstance in which the chosen x-ray mammogram view is a standardized x-ray mammogram view such as the CC or MLO view, the above diagnoses can be quickly and readily made because the screening radiologist already knows and is familiar with the orientations of the ultrasound image slices.

FIG. 5 illustrates a more detailed view of the ultrasound image slice display 406 of the user display of FIG. 4. The medium of the slice display 406 may be any of a variety of display media sufficient to display ultrasound image slice information, such as a TV monitor or a film on a light box. Preferably, the ultrasound image slice display shows a minimum amount of information necessary for the screening radiologist to readily understand the significance of the ultrasound image slices being displayed. Thus, for example, ultrasound image slice display 406 comprises a legend 502 showing the chosen



x-ray mammogram view information (“CC” in the example of FIG. 5), as well as information on the spacing of the ultrasound image slices. The spacing of the ultrasound image slices is of particular importance to the screening radiologist, as it raises awareness of whether a “complete” picture of closely spaced slices is being shown, or whether there is undersampling due to large spaces between the planes.

FIG. 6 illustrates steps taken by a breast cancer screening system in accordance with another preferred embodiment, in which ultrasound data is used to generate ultrasound image slices for display to a screening radiologist, who is familiar with the standard x-ray mammogram views but without the aid of corresponding x-ray mammogram views being shown. At step 602, a standardized x-ray mammogram view plane is chosen, *e.g.*, the CC or MLO view. Preferably, a breast cancer screening system that only shows ultrasound image slice information should show ultrasound image slices corresponding to standardized x-ray mammogram views in order to present familiar orientations to the screening radiologist. Subsequent to step 602, steps 604, 606, 608, and 610 are performed in a manner similar to the steps 106, 108, 110, and 112 of FIG. 1, respectively. Finally, at step 612 the ultrasound image slices for the chosen standard x-ray mammogram view plane are displayed to the screening radiologist.

FIG. 7 illustrates a user display 700 for carrying out step 612 of FIG. 6. User display 700 comprises an ultrasound slice display 706 for simultaneously displaying a plurality of ultrasound image slices  $m_j$  adjacent to one another, and additionally comprises elements 708, 710, and 712 similar to elements 408, 410, and 412 of FIG. 4. The ultrasound slice display 706 further comprises a legend 714 similar to the legend 502 of FIG. 5. The embodiments of FIGS. 6 and 7 are advantageous in that they are lower-cost and less complex than the embodiments of FIGS. 1 and 4, and may be of potential use in very low-cost, integrated, portable ultrasound systems for mass breast cancer screening in impoverished areas of the world. Alternatively, the embodiments of FIGS. 6 and 7 may be of use if advances in ultrasound technology yield ultrasound image slice outputs that are of equal or superior resolution than x-ray mammograms. In general, however, it will usually be preferable to combine the ultrasound image slice outputs with x-ray mammogram outputs in accordance with the embodiments of FIGS. 1 and 4, especially since microcalcifications are not generally detectable using known ultrasound methods, in which

case it will be desirable to obtain and analyze x-ray mammogram data in any case to detect microcalcifications.

FIG. 8 illustrates steps taken by a breast cancer screening system in accordance with another preferred embodiment, in which computer aided diagnosis (CAD) algorithms are performed on the ultrasound image slices and/or the x-ray mammogram view and the results are flexibly displayed to the screening radiologist. For example, in the embodiment of FIG. 8, steps 802-812 are carried out in a manner similar to steps 102-112 of FIG. 1, respectively. Prior to a display step, however, a step 813 for performing computer aided diagnosis (CAD) algorithms on the ultrasound image slices  $m_j$  is carried out. Optionally, a step 815 for performing CAD algorithms on the chosen x-ray mammogram view is carried out. Finally, at step 817 the x-ray mammogram view, the ultrasound image slices, and the ultrasound image slice CAD results and/or the x-ray mammogram view CAD results are displayed to the screening radiologist.

FIG. 9 illustrates a user display 900 for carrying out step 817 of FIG. 8, the user display 900 comprising elements 902-912 similar to elements 402-412 of FIG. 4, respectively. User display 900 also includes, however, a CAD result display 914 for displaying the ultrasound image slice CAD results and/or the x-ray mammogram view CAD results. In accordance with a preferred embodiment, CAD result display 914 displays the ultrasound image slice CAD results until the user presses a toggle control 916, wherein the CAD result display 914 then displays the x-ray mammogram view CAD results, and vice versa. CAD result display 914 is designed to highlight areas of suspicion in the medical image and draw them to the attention of the screening radiologist, who may then more closely examine the original medical image. As a nonlimiting example, FIG. 9 shows highlighting arrows 918 to draw the attention of the radiologist toward the suspicious lesions appearing in the ultrasound image slices.

Any of a variety of ultrasound CAD algorithms can be used at step 813 depending on the desired clinical application including, but not limited to, algorithms disclosed in Chen et. al., "Computer-aided Diagnosis Applied to Ultrasound of Solid Breast Nodules by Using Neural Networks," *Radiology*, pp. 407-412 (November 1999), Cheng et. al., "Automated Detection of Breast Tumors in Ultrasonic Images Using Fuzzy Reasoning," *supra*, and U.S. Pat. 5,984,870 to Giger et. al., *supra*, the contents of each of these

references hereby being incorporated by reference. Also, any of a variety of x-ray mammogram CAD algorithms can be used at step 815 depending on the desired clinical application including, but not limited to, algorithms disclosed in U.S. Pat. 5,815,591 to Roehrig et. al., the contents of which are hereby incorporated by reference.

5           FIG. 10 illustrates steps taken by a breast cancer screening system in accordance with another preferred embodiment in which vibrational resonance information is flexibly displayed to the screening radiologist. At step 1002 the primary ultrasound scan data is obtained. As used herein, primary ultrasound data refers to any type of ultrasound data acquired while the breast is not being acoustically vibrated by an external transducer. The  
10 primary ultrasound data may generally be of any type including B-mode, color Doppler, power Doppler, and other types. At step 1004, a three-dimensional volume reconstruction of the breast is performed using the primary ultrasound scan data. At step 1006, vibrational resonance ultrasound scan data is acquired in a manner similar to that described in Lowers, *supra*, by introducing acoustic vibrations into the breast, allowing the acoustic  
15 vibrations to stabilize, and taking power Doppler ultrasound scans. At step 1008, the power Doppler image slices are computed. At step 1010, a 3-D volume reconstruction of the breast is performed using the vibrational resonance information. As used herein, vibrational resonance information refers to ultrasound scan data taken while the breast is acoustically vibrated; vibrational resonance image slice refers to an ultrasound image slice  
20 derived from the vibrational resonance information; vibrational resonance volume refers to a 3-D volume reconstruction computed from the vibrational resonance image slices; and vibrational resonance image refers to refers to either a vibrational resonance image slices or to an ultrasound image derived from the vibrational resonance volume. The spatial resolution of the vibrational resonance information, and the volumes/images derived  
25 therefrom, will usually be less than the spatial resolution of the primary ultrasound information due to the greater amount of processing required for power Doppler images, although the scope of the preferred embodiments is not so limited.

At step 1012, the choice of x-ray mammogram view plane is received (*e.g.*, CC or MLO) and at step 1014 the corresponding mammogram is obtained. Importantly, the steps  
30 1002, 1006, and 1012 may be carried out in any order and may occur at substantially different times. At steps 1016 and 1018, respectively, primary and vibrational resonance

images are computed from their corresponding 3-D image volumes along planes parallel to the chosen x-ray mammogram plane. At step 1020, the x-ray mammogram and primary ultrasound image slices are displayed to the user using methods described *supra*. At step 1022, to obtain a more thorough view of a given lesion or feature, the user may activate a  
5 simple toggle switch or equivalent keyboard or mouse entry that allows the vibrational resonance images to be superimposed upon the primary ultrasound images. The user may easily toggle between “superimposed” and “non-superimposed” states during the examination. In this way, vibrational resonance information easily and intuitively integrated with the primary ultrasound display in a manner readily usable by the screening  
10 radiologist or other medical professional.

It has been found that different amounts of tissue type differentiation may be acquired for different vibrational frequencies and/or amplitudes. According to a preferred embodiment, “N” distinct sets of vibrational resonance information are acquired for different acoustic vibration frequencies and/or amplitudes. For example, a first set may be  
15 acquired for an acoustic frequency of 100 Hz at 20 watts of acoustic power, a second set for 200 Hz at 20 watts, a third set for 200 Hz at 40 watts, and so on. Each of the “N” data sets yields its own vibrational resonance volume and its own set of vibrational resonance images, which may be separately displayed to the user according to the methods described herein. Moreover, lesion features may be extracted from each of the “N” vibrational  
20 resonance data sets and provided to the CAD classifiers, *e.g.*, artificial neural networks described herein for classifying the lesions. Advantageously, providing “N” different vibrational resonance data sets to the classifier algorithms accommodates the possibility of locating different lesion types that may respond differently to different vibrational frequencies and amplitudes. Thus, it is to be appreciated that while described herein in  
25 terms of only a single set of vibrational resonance data for simplicity and clarity of explanation, the scope of the preferred embodiments includes the acquisition and use of “N” sets of vibrational resonance data in data display and CAD algorithms.

FIG. 11 illustrates steps taken by a breast cancer screening system in accordance with another preferred embodiment, in which computer aided diagnosis (CAD) algorithms  
30 are performed using the primary ultrasound image data, the x-ray mammogram data, and the vibrational resonance information and the results are flexibly displayed to the screening

radiologist. Steps 1102-1114 are carried out in a manner similar to steps 1002-1014 of FIG. 10, respectively. Prior to a display step, however, a step 1116 for performing computer aided diagnosis (CAD) algorithms using the primary ultrasound image data, the x-ray mammogram data, and the vibrational resonance information. Finally, at step 1118  
5 the x-ray mammogram view, the primary ultrasound image slices, the vibrational resonance images, and the CAD results are displayed to the screening radiologist in any of a variety of combinations. In particular, the CAD results may be superimposed on any of the above medical images, or any combination thereof, in accordance with a preferred embodiment. The various steps of FIG. 11 may be carried out in different orders, may  
10 occur at substantially different times, and may be carried out by different, physically separated hardware systems.

FIG. 12A illustrates steps for computer-aided diagnosis in accordance with a preferred embodiment corresponding to step 1116 of FIG. 11. At steps 1202, 1204, and 1206, respectively, the initial data for the x-ray mammogram information, primary  
15 ultrasound information, and vibrational resonance information is read into the CAD system. At steps 1208, 1210, and 1212, respectively, lesions in the patient are separately localized, *i.e.*, areas containing possible lesions are initially identified, using the respective x-ray, primary ultrasound, and vibrational resonance data. At step 1214, lists of localized lesions are compared and adjusted as necessary. Advantageously, the use of vibrational  
20 resonance data may result in localization of lesions that are initially missed by the primary ultrasound localization algorithms because, as discussed *supra*, some tumors may appear isoechoic with surrounding tissue on the primary ultrasound scans, but may appear hyperechoic or hypoechoic in the vibrational resonance images. Most commonly, the list of localized lesions will simply comprise the logical union of lesions localized at steps  
25 1208-1212, although the scope of the preferred embodiments is not so limited.

At steps 1216, 1218, and 1220, respectively, features of the lesions are separately extracted, using the respective x-ray, primary ultrasound, and vibrational resonance data. The various features extracted from the x-ray mammogram and the primary ultrasound data include those discussed in Giger, *supra*, including gray-level based features,  
30 geometric features, gradient features, and other features. Features extracted from the vibrational resonance data include a first set similar to those extracted from the primary

ultrasound data, and a second set based on vibrational resonance parameters. According to a preferred embodiment, this second set of features includes differential gray scale metrics, differential geometric feature metrics, and differential gradient metrics measured versus changes in vibrational resonance frequency and vibrational resonance power. Differential

5 metrics further include differences in a given metric between a vibrational resonance image and the primary ultrasound image (for which the vibrational resonance frequency and vibrational resonance power are both equal to zero).

By way of example, gray-level based features (which for ultrasonic data refers to acoustic impedance amplitude, power Doppler amplitude, etc.) include: average level

10 within a region of interest (ROI) or lesion; standard deviation of the levels within a ROI or lesion; minimum level within a ROI or lesion; maximum level within a ROI or lesion; minimum 5% level corresponding to the level below which yields 5% of the area under the level histogram; maximum 5% level corresponding to the level above which yields 5% of the area under the level histogram; and contrast as given by the difference between levels

15 within the lesion relative to those in the surrounding area. Geometric features include, for example, ratio of lesion surface area to lesion volume.

By way of example, differential gray scale metrics include: a ratio of primary ultrasound image contrast for the lesion versus vibrational resonance image contrast for the lesion; ratio of standard deviation in the lesion for the primary ultrasound image versus

20 standard deviation in the lesion for the vibrational resonance image; first and second derivatives of vibrational resonance image contrast versus vibrational resonance frequency; first and second derivatives of vibrational resonance image contrast versus vibrational resonance power; and any of a variety of metrics that compare (i) the primary image versus a vibrational resonance image, or (ii) different vibrational resonance images at different

25 vibrational resonance frequencies and powers. It is to be appreciated that differential metrics based on the primary image versus the vibrational resonance image will require communication of results between hardware carrying out steps 1218 and 1220.

Differential geometric features include, for example, the surface area/volume ratio for the ultrasound image divided by the surface area/volume ratio for the vibrational resonance

30 image.

At step 1222, classifier algorithms, *e.g.* artificial neural network algorithms, are executed on the features extracted at steps 1216-1220, and at step 1224 a degree of user attention required for that feature (*e.g.* in the form of a highlighting symbol type, symbol brightness, etc.) is determined. Finally, at step 1226 the CAD results are output for display  
5 in accordance with step 1118 of FIG. 11, *supra*.

FIG. 12B illustrates steps for computer-aided diagnosis in accordance with an alternative preferred embodiment corresponding to step 1116 of FIG. 11, wherein CAD algorithms on the x-ray mammogram information are carried out independently of the ultrasound/vibrational resonance CAD algorithms. As indicated in FIG. 12B, steps 1202-  
10 1212 are carried out in the same manner as for FIG. 12A. However, beginning at step 1214A, the x-ray mammogram CAD algorithm steps are carried out independently, while the ultrasound/vibrational resonance CAD algorithms are still executed together. Thus, at step 1214A, only lists of localized lesions from the ultrasound and vibrational resonance images are compared and adjusted as necessary. Feature extraction step 1216A,  
15 classification step 1222A, user attention determining step 1224A are separately executed on the x-ray mammogram data without using any data from the ultrasound/vibrational resonance CAD algorithms. At step 1226A, the x-ray mammogram CAD results are output separately for display to the user. Likewise steps 1218A, 1220A, and 1222B-1226B are carried out without using x-ray mammogram data or CAD results. Accordingly, in the  
20 preferred embodiment of FIG. 12B, the first integration between the x-ray mammogram CAD results and the ultrasound/vibrational resonance CAD results is performed by the user when simultaneously viewing the respective displays.

FIG. 13 illustrates a breast ultrasound adapter 1300 for facilitating breast ultrasound scans in accordance with a preferred embodiment. As known in the art, it is  
25 required that a close acoustical coupling be maintained between the breast skin surface and the ultrasound probe at all times during the scan. In particular, there should be no vacuum or air pockets anywhere along the path of the interrogating ultrasound pulses. This is usually achieved by keeping the breast surface wet with a water-containing polymeric fluid or gel at all points of contact with the ultrasound probe. Alternatively, gel bags have been  
30 placed between the ultrasound probe and skin surface to maintain an acoustic coupling.

With both of these methods, however, the breast itself is necessarily shifted around and compressed by differing amounts at differing places as the ultrasound probe is moved.

Breast ultrasound adapter 1300 comprises an upper frame 1302, side walls 1306, and a bottom membrane 1310 forming a fluid reservoir volume 1308 for holding a water-  
5 containing polymeric fluid or gel. As shown in FIG. 13, breast ultrasound adapter 1300 has been placed over a breast 1320 and filled with fluid 1311, and the bottom membrane 1310 has conformally adapted to the shape of the breast 1320. The fluid level should rise above the highest point of the breast 1320. The bottom of an ultrasound probe 1314 is submerged in the fluid 1311 and is swept in a lateral direction across the breast as indicated  
10 in FIG. 13, the bottom of the ultrasound probe 1314 remaining submerged during the lateral sweep. Preferably, the bottom of ultrasound probe 1314 is maintained in a plane parallel to the fluid surface and does not contact the bottom membrane 1310 during the lateral sweep. Position sensors (not shown) are used to track the position of the ultrasound probe 1314 as it is swept, usually by hand, across the fluid surface. Preferably, water-  
15 containing polymeric gel, water, and/or oil are used to improve the contact between the bottom membrane 1310 and the breast 1320. Advantageously, the breast is maintained in a substantially consistent position and in a consistent state of compression during the scanning process, thereby facilitating consistency among the ultrasound slices forming the three-dimensional image volume.

20 The breast ultrasound adapter 1300 may comprise any of a variety of material configurations that facilitate the presence of an open fluid reservoir above the breast of a supine patient, the fluid reservoir having a lower surface that conforms to the breast shape such that an ultrasound probe is acoustically coupled to the breast skin surface when immersed in the fluid. By way of example, upper frame 1302 is preferably a rigid or semi-  
25 rigid compression-molded silicone rubber material, or an equivalent material, such that the breast ultrasound adapter 1300 can be supported and moved by manipulating the upper frame 1302 even when full of fluid. Side walls 1306 may also be rigid or semi-rigid. In the embodiment of FIG. 13, side walls 1306 are semi-rigid and sufficiently flexible such that they are conformal with the patient's skin surface along points of intersection 1304  
30 therewith. However, in alternative preferred embodiments, the side walls 1306 do not contact the skin surface directly and therefore may be rigid. In this case, the bottom



membrane protrudes downward from the side walls and hangs down like a plastic bag over the breast when filled with fluid. In still other preferred embodiments, the side walls 1306 are not present at all, and the bottom membrane hangs down directly from the upper frame 1302 over the breast when filled with fluid. In preferred embodiments in which the side  
5 walls 1306 do not contact the skin surface directly or are not present, the breast ultrasound adapter 1300 is supported by the upper frame 1302 and laid gently over the breast, which does not support its entire weight. The bottom membrane 1310 preferably comprises a flexible, watertight, conforming material. An at least partially distensible characteristic assists in ensuring bubble-free contact with the breast skin. Suitable materials include  
10 protective latex, synthetic elastomers, cellophane, or other protective sheath-type materials described in U.S. Pat. 6,039,694, which is incorporated by reference herein. The respective pieces are fastened together by a suitable adhesive such as silicone RTV.

FIG. 14 illustrates a breast ultrasound adapter 1402 having additional features in accordance with a preferred embodiment that, in conjunction with a mechanical probe  
15 translation assembly, forms a semi-automatic breast ultrasound scanning device 1400. Breast ultrasound adapter 1402 comprises an integrated audio speaker 1410 coupled to an electrical source 1412 for use in vibrational resonance applications. The apparatus of FIG. 14 also comprises a conceptual diagram of an ultrasound probe 1404 as it is guided by a mechanical translator 1406, for example by means of a slot 1408. Advantageously, the  
20 probe position may be sensed by mechanical transducers instead of position sensors as required in the embodiment of FIG. 13, *supra*. The mechanical translator may be affixed to the frame of the breast ultrasound adapter 1402, or may alternatively be externally supported.

Finally, breast ultrasound adapter 1402 further comprises a fluid transport opening  
25 and fluid conduit 1414 for coupling to a gravity-feed reservoir (not shown) that is functionally similar to an intravenous (IV) fluid container. In accordance with a preferred embodiment, the gravity-feed reservoir is raised, either by hand or by an automated mechanical assembly, to introduce fluid into the breast ultrasound adapter 1402 after it has been placed on a patient's breast. Following the scan procedure, the gravity-feed reservoir  
30 is lowered to cause fluid to drain out, thereby emptying the breast ultrasound adapter 1402.

The vertical position of the gravity-feed reservoir may also be adjusted used to regulate the level of the fluid during the scan.

Whereas many alterations and modifications of the present invention will no doubt become apparent to a person of ordinary skill in the art after having read the foregoing  
5 description, it is to be understood that the particular embodiments shown and described by way of illustration are in no way intended to be considered limiting. By way of example, while vibrational resonance images are described *supra* as comprising power Doppler images, in other preferred embodiments the vibrational resonance images may comprise color Doppler images or other ultrasound image types taken while the breast is being  
10 vibrated. By way of further example, while the breast ultrasound adapter described *supra* covers a single breast, in other preferred embodiments it is configured to cover both breasts simultaneously, thereby further expediting the ultrasound scanning process. Therefore, reference to the details of the preferred embodiments are not intended to limit their scope, which is limited only by the scope of the claims set forth below.

15

## CLAIMS

What is claimed is:

1. A method for facilitating the detection of breast lesions, comprising the steps of:  
5 displaying an x-ray mammogram view of a breast captured in an x-ray mammogram view plane; and  
displaying near said x-ray mammogram view a plurality of ultrasound image slices, said ultrasound image slices being derived from ultrasound scans of the breast, said ultrasound image slices corresponding to a succession of planes in the breast substantially  
10 parallel to said x-ray mammogram view plane, and said mammogram view and ultrasound image slices being selectively displayed concurrently;  
whereby detection of breast lesions is facilitated by simultaneous viewing of said x-ray mammogram view and the succession of ultrasound image slices corresponding thereto.  
15
2. The method of claim 1, wherein the x-ray mammogram view plane is a standardized x-ray mammogram view plane and said ultrasound image slices correspond to planes in the breast substantially parallel to said standardized x-ray mammogram view plane, whereby a user who is accustomed to analyzing x-ray mammograms captured in  
20 said standardized x-ray mammogram view plane can quickly and readily perceive the orientation and medical significance of said ultrasound image slices alongside said x-ray mammogram view.
3. The method of claim 2, wherein said standardized x-ray mammogram view plane  
25 corresponds to a craniocaudal (CC) view.
4. The method of claim 2, wherein said standardized x-ray mammogram view plane corresponds to a mediolateral oblique (MLO) view.
- 30 5. The method of claim 2, wherein said ultrasound image slices correspond to planes in the breast that are separated by a distance no greater than a predetermined spacing, said

predetermined spacing being selected to reduce the probability of missed lesions due to undersampling along an axis perpendicular to said standardized x-ray mammogram view plane.

5 6. The method of claim 5, wherein said predetermined spacing is no greater than 0.5 cm.

7. The method of claim 2, further comprising the step of obtaining an ultrasound scan of the breast, said step of obtaining an ultrasound scan of the breast comprising the steps  
10 of:

flattening the breast against a plate that is substantially parallel to the standardized x-ray mammogram view plane; and

using an ultrasound probe to transmit ultrasonic waves into the breast and receive reflections therefrom to produce ultrasound scan results.

15

8. The method of claim 7, wherein said ultrasound probe is oriented in a direction substantially perpendicular to said standardized x-ray mammogram view plane.

9. The method of claim 8, further comprising the steps of:  
20 constructing a 3-dimensional matrix from said ultrasound scan results, said 3-dimensional matrix representing an acoustic impedance map of the breast volume; and  
deriving said ultrasound image slices by interpolating portions of said 3-dimensional matrix along planes substantially parallel to said standardized x-ray mammogram plane.

25

10. The method of claim 7, wherein said ultrasound probe is oriented in a direction parallel to said standardized x-ray mammogram view plane, whereby said ultrasound image slices are derived directly from said ultrasound scan results without requiring a 3-dimensional volume reconstruction process.

30

11. The method of claim 2, further comprising the steps of:  
performing computer-aided diagnosis on each ultrasound image slice for detecting suspicious locations therein;  
highlighting said suspicious locations on each ultrasound image slice to produce an  
5 ultrasound image slice computer-aided diagnosis (CAD) display; and  
displaying said ultrasound image slice CAD display on a first display screen;  
wherein said first display screen is placed sufficiently close to said x-ray  
mammogram view for a user to simultaneously view the ultrasound image slice CAD  
display and the x-ray mammogram view.
- 10
12. The method of claim 11, further comprising the steps of:  
performing computer-aided diagnosis on the x-ray mammogram view for detecting suspicious locations therein;  
highlighting said suspicious locations to produce an x-ray mammogram CAD  
15 display;  
receiving a first toggle command from the user;  
responsive to said first toggle command, displaying said x-ray mammogram CAD  
on said first display screen;  
receiving a second toggle command from the user;  
20 responsive to said second toggle command, displaying said ultrasound image slice  
CAD display on said first display screen;  
whereby the user may easily and quickly toggle between said x-ray mammogram  
CAD display and said ultrasound image slice CAD display on said first display screen  
while simultaneously viewing the x-ray mammogram view for increased reliability of  
25 lesion detection in the breast without substantial reduction in speed of analysis.
13. An apparatus for assisting in breast cancer screening, comprising:  
a first display device adapted to display an x-ray mammogram view of a breast  
captured in an x-ray mammogram view plane;  
30 a processor configured to receive ultrasound data derived from ultrasound scans of  
the breast and generate therefrom a plurality of ultrasound image slices, said ultrasound

image slices corresponding to planes in the breast substantially parallel to the x-ray mammogram view plane; and

a second display device configured to display said plurality of ultrasound image slices, wherein said second display is placed sufficiently adjacent to said first display

5 device for simultaneous viewing by a user, whereby said apparatus facilitates detection of breast lesions by convenient simultaneous viewing of individual planes of the breast on said second display device alongside said x-ray mammogram view on said first display device.

10 14. The apparatus of claim 13, the ultrasound scans being taken of the breast when flattened along a plane substantially parallel to the x-ray mammogram view plane, said processor being configured to perform 3-dimensional reconstruction of said ultrasound image slices from said ultrasound data, said 3-dimensional reconstruction comprising the steps of:

15 constructing a 3-dimensional acoustic impedance matrix from said ultrasound data; and

deriving said ultrasound image slices by interpolating portions of said 3-dimensional acoustic impedance matrix along planes substantially parallel to the x-ray mammogram view plane.

20

15. The apparatus of claim 14, the ultrasound scans being taken by an ultrasound probe, said processor being configured to receive a probe orientation signal, said processor being configured to perform said step of 3-dimensional reconstruction of said ultrasound image slices if said probe orientation signal indicates that said probe has scanned along a  
25 plane not substantially parallel to the x-ray mammogram view plane, said processor being configured to directly derive said ultrasound image slices from said ultrasound data without performing 3-dimensional reconstruction if said probe orientation signal indicates that said probe has scanned along a plane substantially parallel to said x-ray mammogram view plane.

30

16. The apparatus of claim 15, wherein the x-ray mammogram view plane is a standardized x-ray mammogram view plane and said ultrasound image slices correspond to planes in the breast substantially parallel to said standardized x-ray mammogram view plane, whereby an experienced radiologist familiar with analyzing x-ray mammograms  
5 captured in said standardized x-ray mammogram view plane can quickly and readily perceive the orientation and medical significance of said ultrasound image slices in analyzing the breast.
17. The apparatus of claim 16, wherein said standardized x-ray mammogram view  
10 plane corresponds to a craniocaudal (CC) view.
18. The apparatus of claim 16, wherein said standardized x-ray mammogram view plane corresponds to a mediolateral oblique (MLO) view.
- 15 19. The apparatus of claim 13, said processor being a first processor, said apparatus further comprising a second processor configured to receive said ultrasound image slices and to perform computer-aided diagnosis (CAD) algorithms thereon for detecting suspicious locations in the breast, wherein said second display device is configured to superimpose highlighting marks at said suspicious locations of said ultrasound image  
20 slices.
20. The apparatus of claim 19, further comprising:  
a third processor configured to receive information from said x-ray mammogram view and to perform computer-aided diagnosis (CAD) algorithms thereon for detecting  
25 suspicious locations in the breast, wherein said second display device is also configured to display a digital representation of said x-ray mammogram view having highlighting marks at said suspicious locations.
21. The apparatus of claim 20, further comprising a toggle switch readily accessible by  
30 the user, wherein the second display device is configured to toggle between the highlighted

ultrasound CAD display and the highlighted x-ray mammogram CAD display responsive to user control of said toggle switch.

22. A computer program product for assisting in breast cancer screening through  
5 computerized manipulation and display of breast images, comprising:  
computer code for causing the display of an x-ray mammogram view of a breast on a first display device, the x-ray mammogram view being captured in an x-ray mammogram view plane;  
computer code for receiving ultrasound data derived from ultrasound scans of the  
10 breast and for generating therefrom a plurality of ultrasound image slices, said ultrasound image slices corresponding to planes in the breast substantially parallel to the x-ray mammogram view plane; and  
computer code for causing the display of said ultrasound image slices on a second display device, the second display device being sufficiently near said first display device  
15 for simultaneous viewing of the first display device and the second display device by a user.

23. The computer program product of claim 22, wherein said first display device is a mechanical display apparatus for displaying x-ray mammograms recorded on x-ray film,  
20 and wherein said second display device comprises a computer monitor.

24. The computer program product of claim 22, wherein the x-ray mammogram view plane is a standardized x-ray mammogram view plane and said ultrasound image slices correspond to planes in the breast substantially parallel to said standardized x-ray  
25 mammogram view plane, whereby an experienced radiologist familiar with analyzing x-ray mammograms captured in said standardized x-ray mammogram view plane can quickly and readily perceive the orientation and medical significance of said ultrasound image slices in analyzing the breast.

- 30 25. The computer program product of claim 24, wherein said standardized x-ray mammogram view plane corresponds to a craniocaudal (CC) view.



26. The computer program product of claim 24, wherein said standardized x-ray mammogram view plane corresponds to a mediolateral oblique (MLO) view.

5 27. The computer program product of claim 24, the ultrasound scans being taken of the breast when flattened along a plane substantially parallel to the x-ray mammogram view plane, further comprising computer code for performing 3-dimensional reconstruction of said ultrasound image slices from said ultrasound data, wherein said computer code for performing 3-dimensional reconstruction comprises:

10 computer code for constructing a 3-dimensional acoustic impedance matrix from said ultrasound data; and

computer code for deriving said ultrasound image slices by interpolating portions of said 3-dimensional acoustic impedance matrix along planes substantially parallel to the x-ray mammogram view plane.

15

28. The computer program product of claim 27, the ultrasound scans being taken by an ultrasound probe, further comprising:

computer code for receiving a probe orientation signal indicating an orientation of the ultrasound probe with respect to the x-ray mammogram view plane;

20 computer code for invoking said computer code for performing 3-dimensional reconstruction of said ultrasound image slices if said probe orientation signal indicates that said ultrasound probe has scanned along a plane not substantially parallel to the x-ray mammogram view plane; and

computer code for directly deriving said ultrasound image slices from said  
25 ultrasound data without performing 3-dimensional reconstruction if said probe orientation signal indicates that said probe has scanned along a plane substantially parallel to said x-ray mammogram view plane.

29. The computer program product of claim 28, further comprising:

computer code for receiving said ultrasound image slices and performing computer-aided diagnosis (CAD) algorithms thereon for detecting suspicious locations in the breast, and

5 computer code for highlighting said second display device at said suspicious locations of said ultrasound image slices.

30. The computer program product of claim 29, further comprising:

computer code for receiving information from said x-ray mammogram view and performing computer-aided diagnosis (CAD) algorithms thereon for detecting suspicious  
10 locations in the breast;

computer code for causing said second display device to display a digital representation of said x-ray mammogram view having highlighting marks at said suspicious locations.

15 31. The computer program product of claim 30, further comprising:

computer code for receiving toggle commands from the user; and  
computer code for causing said second display device to toggle between said highlighted ultrasound CAD display and said highlighted x-ray mammogram CAD display responsive to said toggle commands.

20

32. A method for facilitating the detection of breast lesions, comprising the steps of:

obtaining ultrasound data from an ultrasound scan of a breast;  
computing from said ultrasound data a plurality of ultrasound image slices, said ultrasound image slices corresponding to a succession of parallel planes in the breast; and

25 displaying said plurality of ultrasound image slices on a user display;

wherein said ultrasound image slices correspond to planes in the breast substantially parallel to a standardized x-ray mammogram view plane;

whereby a user accustomed to analyzing x-ray mammograms captured in said standardized x-ray mammogram view plane can quickly and readily perceive the  
30 orientation and medical significance of said ultrasound image slices in analyzing the breast.

33. The method of claim 32, wherein said standardized x-ray mammogram view plane is selected from a group of standardized x-ray mammogram view planes, said group comprising a mediolateral oblique (MLO) view and a craniocaudal (CC) view.

5

34. The method of claim 33, further comprising the steps of:  
performing computer-aided diagnosis on each ultrasound image slice for detecting suspicious locations therein;

10 highlighting said suspicious locations to produce an ultrasound image slice  
computer-aided diagnosis (CAD) display; and  
displaying said ultrasound image slice CAD display near said plurality of  
ultrasound image slices on said user display.

35. The method of claim 33, wherein said succession of ultrasound image slices  
15 correspond to planes in the breast that are separated by a distance no greater than a  
predetermined spacing, said predetermined spacing being selected to reduce the probability  
of missed lesions due to undersampling along an axis perpendicular to said standardized x-  
ray mammogram view plane.

20 36. The method of claim 35, wherein said predetermined spacing is no greater than 0.5  
cm.

37. The method of claim 32, further comprising the step of obtaining an ultrasound  
scan of the breast, said step of obtaining an ultrasound scan of the breast comprising the  
25 steps of:

flattening the breast against a plate that is substantially parallel to the standardized  
x-ray mammogram view plane; and

using an ultrasound probe to transmit ultrasonic waves into the breast and receive  
reflections therefrom to produce ultrasound scan results.

30

38. The method of claim 37, wherein said ultrasound probe is oriented in a direction substantially perpendicular to said standardized x-ray mammogram view plane.

39. The method of claim 38, further comprising the steps of:

- 5 constructing a 3-dimensional matrix from said ultrasound scan results, said 3-dimensional matrix representing an acoustic impedance map of the breast volume; and deriving said ultrasound image slices by interpolating portions of said 3-dimensional matrix along planes substantially parallel to said standardized x-ray mammogram plane.

10

40. The method of claim 38, wherein said ultrasound probe is oriented in a direction parallel to said standardized x-ray mammogram view plane, whereby said ultrasound image slices are derived directly from said ultrasound scan results without requiring a 3-dimensional volume reconstruction process.

15

41. A method for facilitating the detection of breast lesions, comprising the steps of: displaying an x-ray mammogram view of a breast captured in an x-ray mammogram view plane;

- 20 displaying near said x-ray mammogram view a plurality of ultrasound image slices, said ultrasound image slices being derived from a first set of ultrasound scans of the breast, said ultrasound image slices corresponding to a succession of planes in the breast substantially parallel to said x-ray mammogram view plane; and

25 displaying near said ultrasound image slices a corresponding plurality of vibrational resonance image slices derived from a second set of ultrasound scans of the breast, said second set of ultrasound scans being taken while the breast is being vibrated at an audio frequency;

whereby detection of breast lesions is facilitated by simultaneous viewing of said x-ray mammogram view, said succession of ultrasound image slices, and said corresponding vibrational resonance image slices.

30

42. The method of claim 41, wherein said vibrational resonance images are superimposed upon said ultrasound image slices.
43. The method of claim 41, further comprising:  
5 receiving a first toggle command from a user;  
responsive to receiving said first toggle command, superimposing at least one of said vibrational resonance images upon at least one of said ultrasound image slices;  
receiving a second toggle command from the user; and  
responsive to receiving said second toggle command, removing said vibrational  
10 resonance image from said ultrasound image slice.
44. The method of claim 41, wherein said second set of ultrasound scans from which said vibrational resonance images are derived comprise power Doppler image information.
- 15 45. The method of claim 41, further comprising:  
performing computer-assisted detection (CAD) algorithms using information from said x-ray mammogram view, said ultrasound image slices, and said vibrational resonance image slices for detecting one or more suspicious lesions in the breast; and  
highlighting at least one of said x-ray mammogram, ultrasound slice, or vibrational  
20 resonance image displays at locations corresponding to said suspicious lesions.
46. The method of claim 45, wherein performing CAD algorithms comprises:  
localizing a candidate set of lesions using information from said x-ray mammogram view, said ultrasound image slices, and said vibrational resonance images;  
25 performing feature extraction algorithms on said candidate lesions using information from said x-ray mammogram view, said ultrasound image slices, and said vibrational resonance images; and  
performing classifier algorithms on each candidate lesion using at least one feature extracted from each of said x-ray mammogram view, said ultrasound image slices, and said  
30 vibrational resonance images.

47. The method of claim 46, wherein one of said extracted features is computed by dividing a lesion contrast metric from the ultrasound image slices by a lesion contrast metric from the vibrational resonance images.
- 5 48. The method of claim 46, wherein "N" sets of vibrational resonance images are acquired for different audio frequencies and/or powers, and wherein one of said extracted features includes a differential vibrational resonance feature computed using two or more of said "N" sets of vibrational resonance images.
- 10 49. The method of claim 48, wherein said differential vibrational resonance feature is computed based on a difference between (i) a first lesion contrast metric computed from a first set of vibrational resonance images taken at a first audio frequency, and (ii) a second lesion contrast metric computed from a second set of vibrational resonance images taken at a second audio frequency.
- 15 50. The method of claim 48, wherein said differential vibrational resonance feature is computed based on a difference between (i) a first surface area/volume metric computed from a first set of vibrational resonance images taken at a first audio frequency, and (ii) a second surface area/volume metric computed from a second set of vibrational resonance
- 20 images taken at a second audio frequency.
51. A computer program product for identifying suspicious lesions in a breast, comprising:
- 25 computer code for receiving x-ray mammogram information corresponding to the breast;
- computer code for receiving vibrational resonance image information derived from a first ultrasound scan of the breast acquired as the breast was vibrated at an audio frequency;
- computer code for receiving primary ultrasound image information derived from a
- 30 second ultrasound scan of the breast acquired when the breast was non-vibrated;

computer code for localizing a candidate set of lesions using said x-ray mammogram information, said vibrational resonance image information, and said primary ultrasound image information;

5 computer code for extracting features of each of said candidate lesions using said x-ray mammogram information, said vibrational resonance image information, and said primary ultrasound image information; and

computer code for classifying each of said candidate lesions using at least one of said extracted features.

10 52. The computer program product of claim 51, wherein one of said extracted feature is computed by dividing a lesion contrast metric from the ultrasound image information by a lesion contrast metric from the vibrational resonance information.

53. The computer program product of claim 51, further comprising:

15 computer code for receiving "N-1" additional sets of vibrational resonance image information derived from "N-1" additional ultrasound scans of the breast acquired as the breast was vibrated at "N-1" different audio frequencies or audio powers; and

computer code for extracting a differential vibrational resonance feature computed using at least two sets of said "N" sets of vibrational resonance images.

20

54. The computer program product of claim 53, wherein said differential vibrational resonance feature is computed based on a difference between (i) a first lesion contrast metric computed from a first of said "N" sets of vibrational resonance images taken at a first audio frequency, and (ii) a second lesion contrast metric computed from a second of  
25 said "N" sets of vibrational resonance images taken at a second audio frequency.

55. The computer program product of claim 53, wherein said differential vibrational resonance feature is computed based on a difference between (i) a first surface area/volume ratio computed from a first of said "N" sets of vibrational resonance images taken at a first  
30 audio frequency, and (ii) a second surface area/volume ratio computed from a second of said "N" sets of vibrational resonance images taken at a second audio frequency.

56. An apparatus for acoustically coupling an ultrasound probe to a breast of a supine patient, comprising:

an upper rim defining an entrance to an open fluid reservoir, said entrance being  
5 large enough to allow the ultrasound probe to pass into the reservoir to achieve contact with fluid being held therein; and

a flexible membrane defining a lower boundary of the reservoir, said flexible membrane coming into conforming contact with the breast surface when the reservoir is aligned with the breast and is at least partially filled with fluid, a substantially flat fluid  
10 surface being achieved when the upper rim is placed above a highest breast point and the fluid level is raised above said highest breast point, an acoustic coupling between the ultrasound probe and the breast being achieved when the ultrasound probe is at least partially submerged in the fluid.

15 57. The apparatus of claim 56, said flexible membrane comprising a distensible, biocompatible, acoustically conducting, waterproof material.

58. The apparatus of claim 56, said entrance defined by said upper rim having a cross-sectional area roughly corresponding to a lateral extent of the breast of the supine patient.

20

59. The apparatus of claim 56, said entrance defined by said upper rim having a cross-sectional area roughly corresponding to a lateral extent of the both breasts of the supine patient, said flexible membrane being sufficiently large to come into conforming contact with both breast surfaces when the reservoir is aligned with the breasts and is at least  
25 partially filled with fluid.

60. The apparatus of claim 56, said upper rim being substantially rigid, said apparatus further comprising side walls fastened at an upper edge to said upper frame and fastened near a lower edge to said flexible membrane, said side walls being substantially rigid, said  
30 flexible membrane extending beneath said side walls when said reservoir is filled with fluid for coming into conformal contact with the breast surface.



61. The apparatus of claim 56, said upper rim being substantially rigid, said apparatus further comprising side walls fastened at an upper edge to said upper frame and fastened at a lower edge to said flexible membrane, said side walls being sufficiently flexible such that  
5 said side walls also conform to the patient when said reservoir is lowered onto the breast and filled with fluid.

62. The apparatus of claim 56, further comprising an audio transducer for acoustically vibrating the breast at one or more audio frequencies during a vibrational resonance  
10 ultrasound procedure, said audio transducer being positioned to acoustically communicate with the reservoir fluid when said reservoir is filled past a predetermined level.

63. The apparatus of claim 56, further comprising an external, vertically movable gravity-feed fluid container, said container being in fluid communication with said  
15 reservoir such that fluid is transferred from the container to the reservoir when a fluid level of the container is raised above a fluid level of the reservoir, and such that fluid is emptied from the reservoir to the container when the fluid level of the container is lowered below the fluid level of the reservoir.

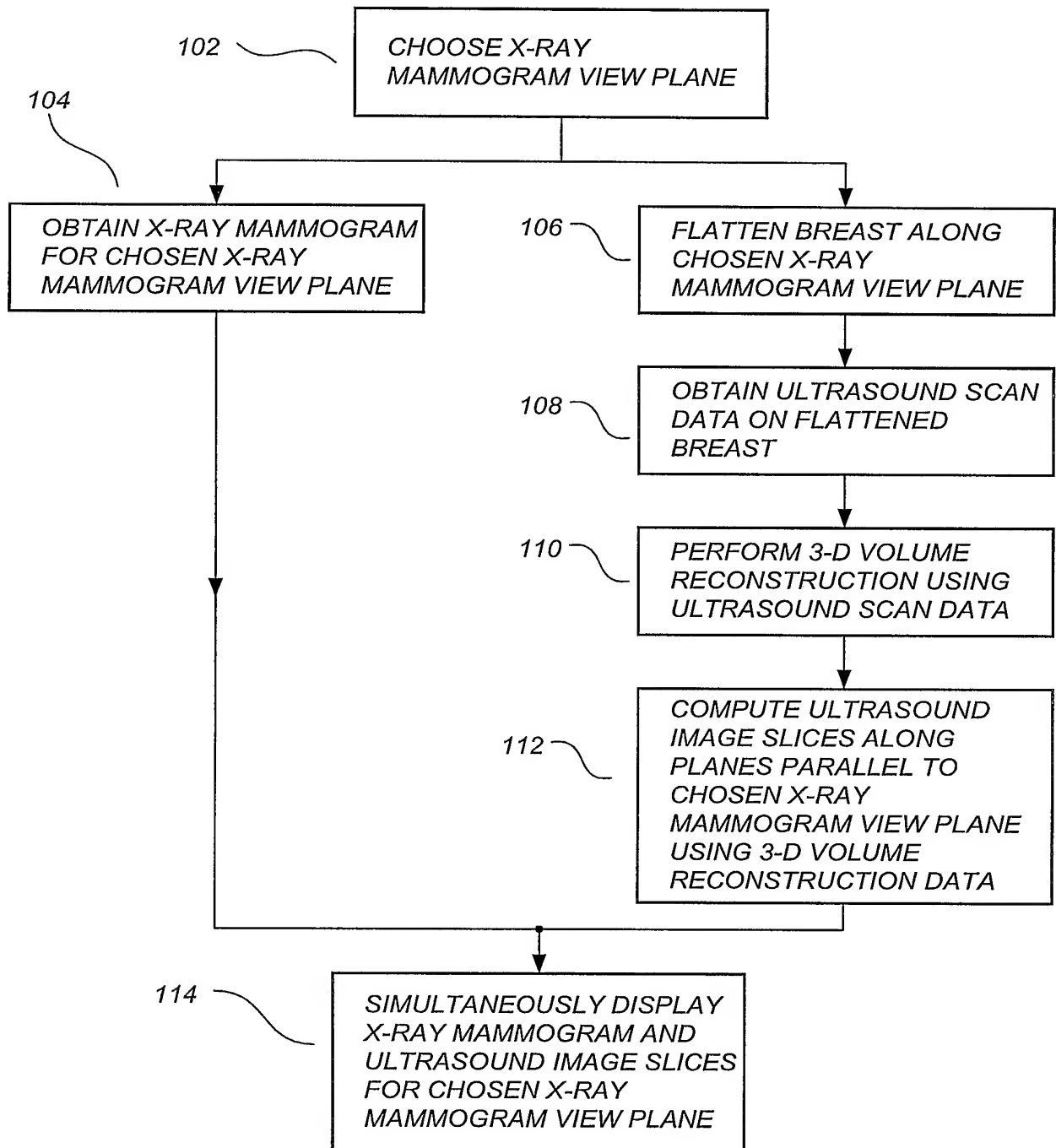
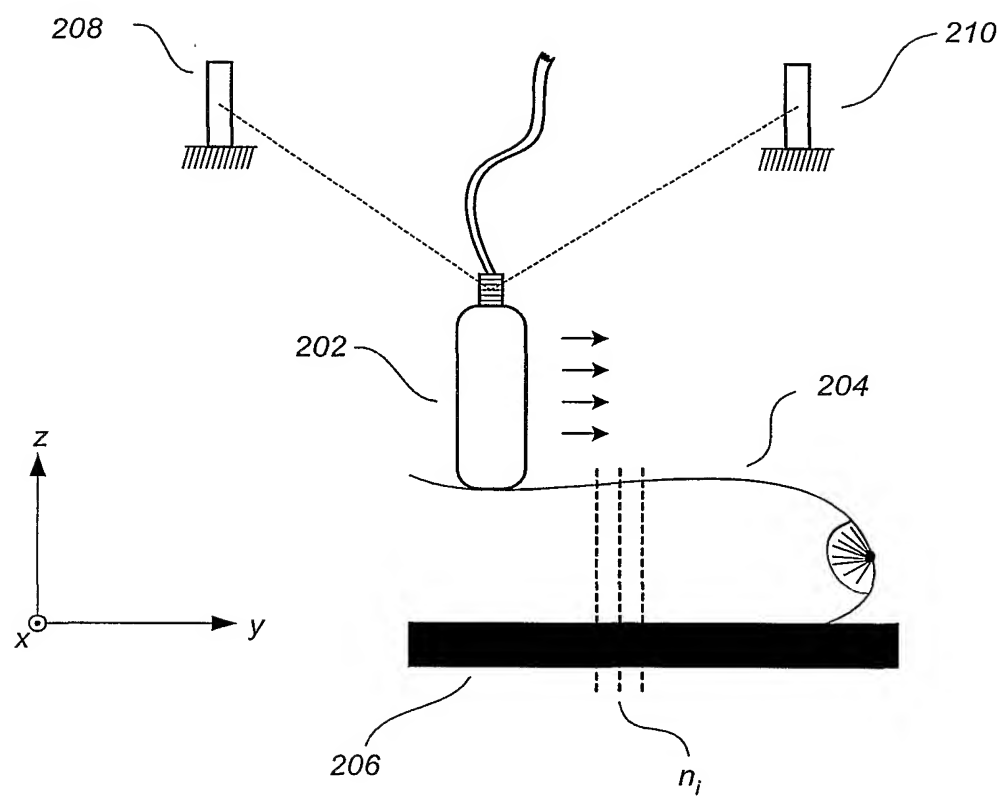


FIG. 1

**FIG. 2**

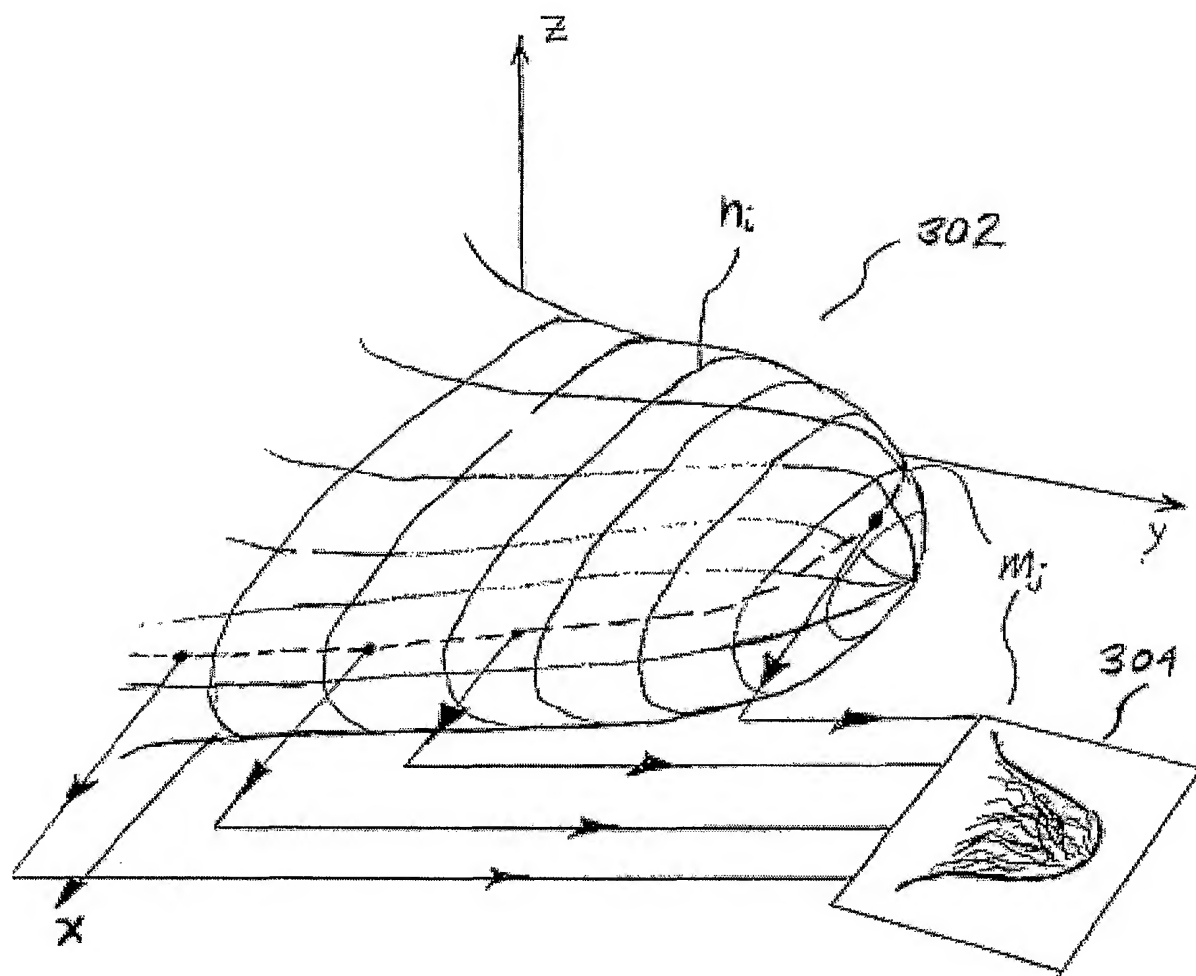


FIG. 3

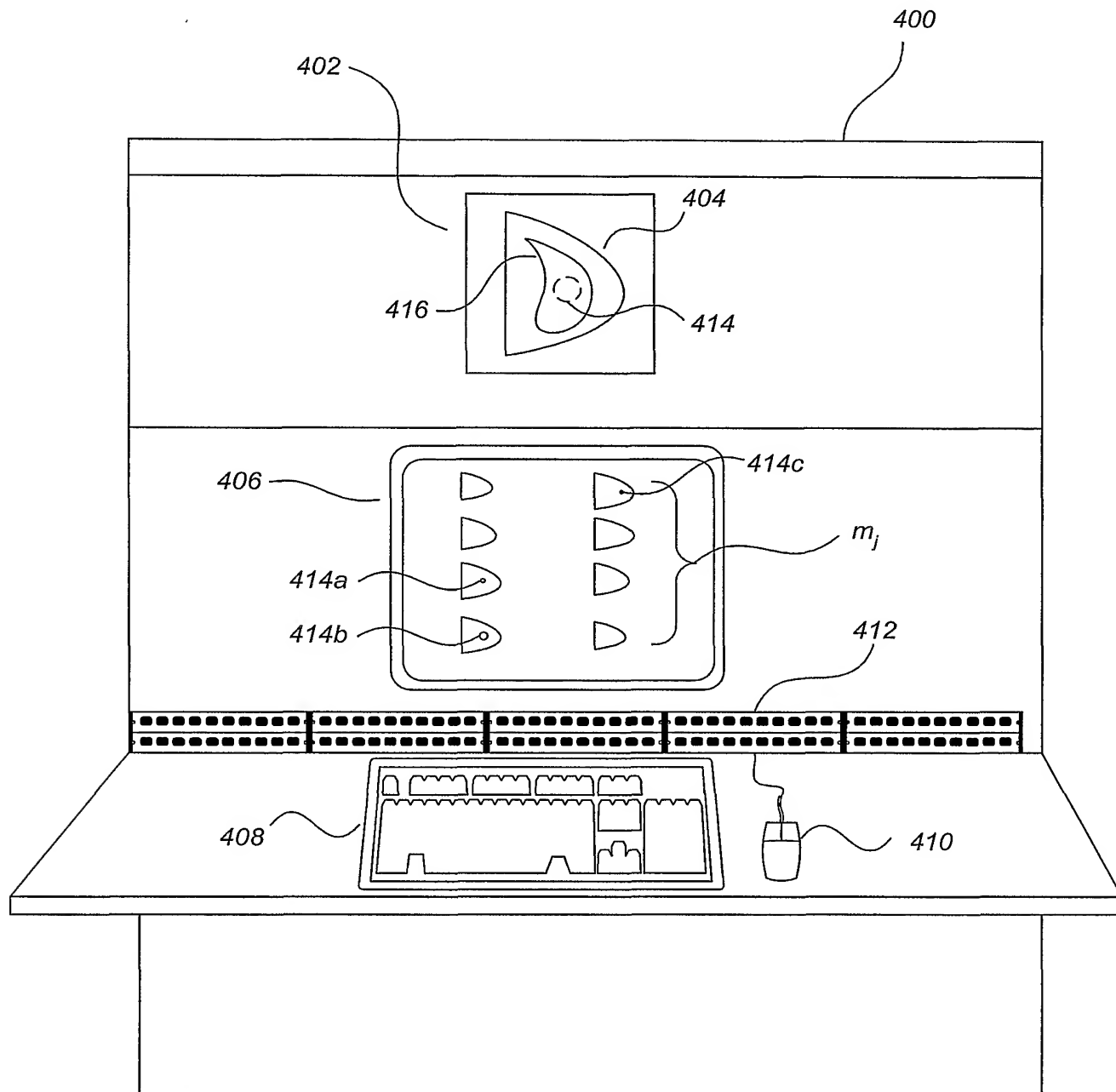


FIG. 4

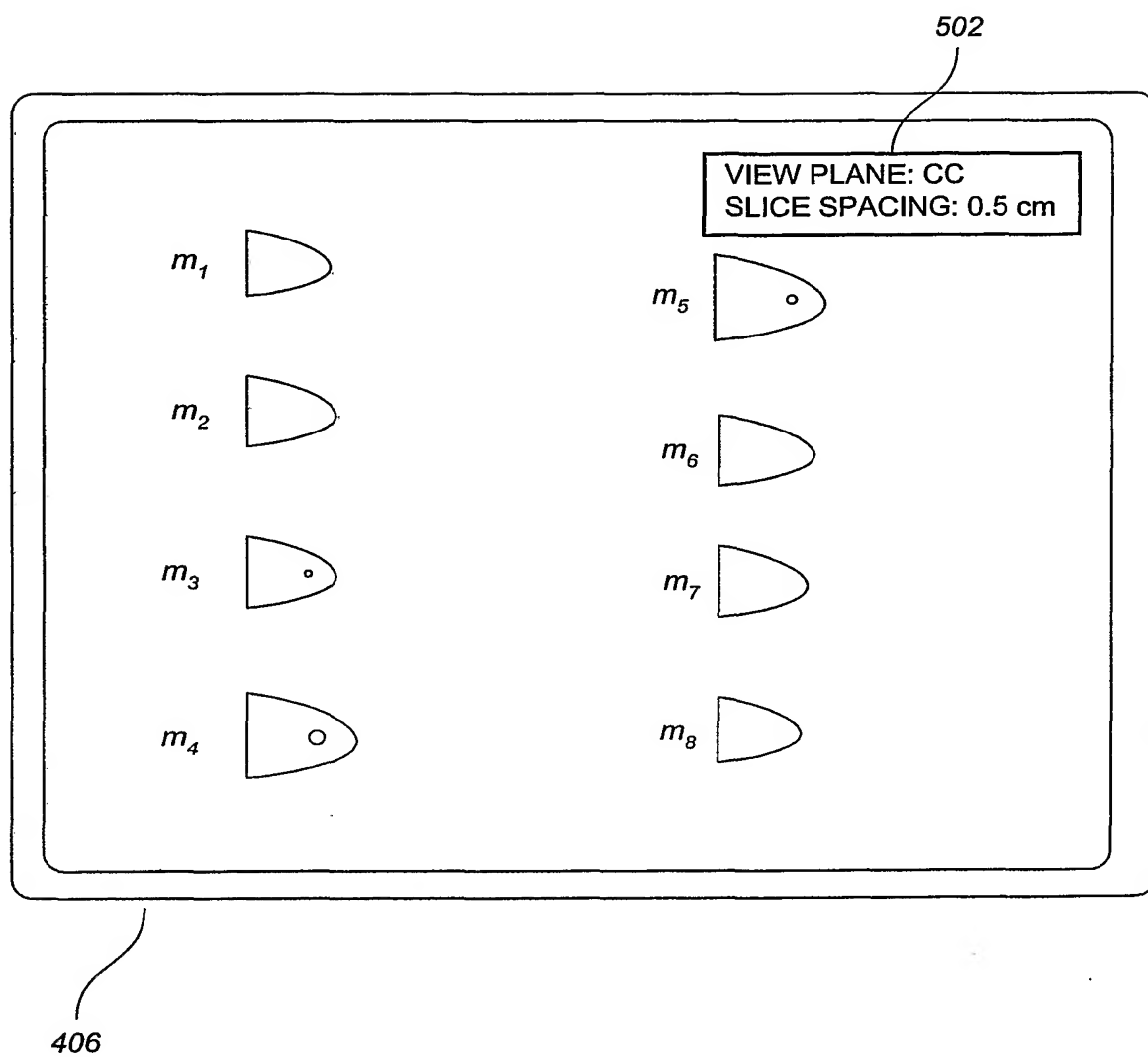
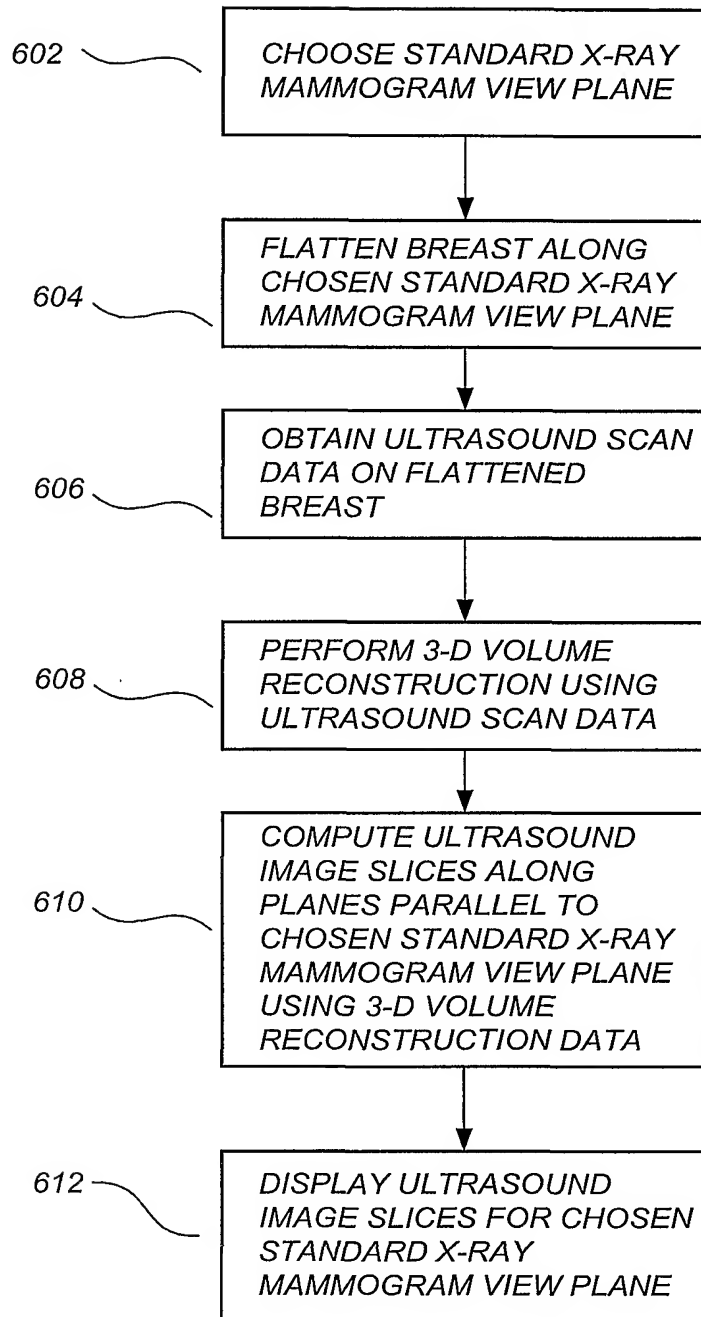
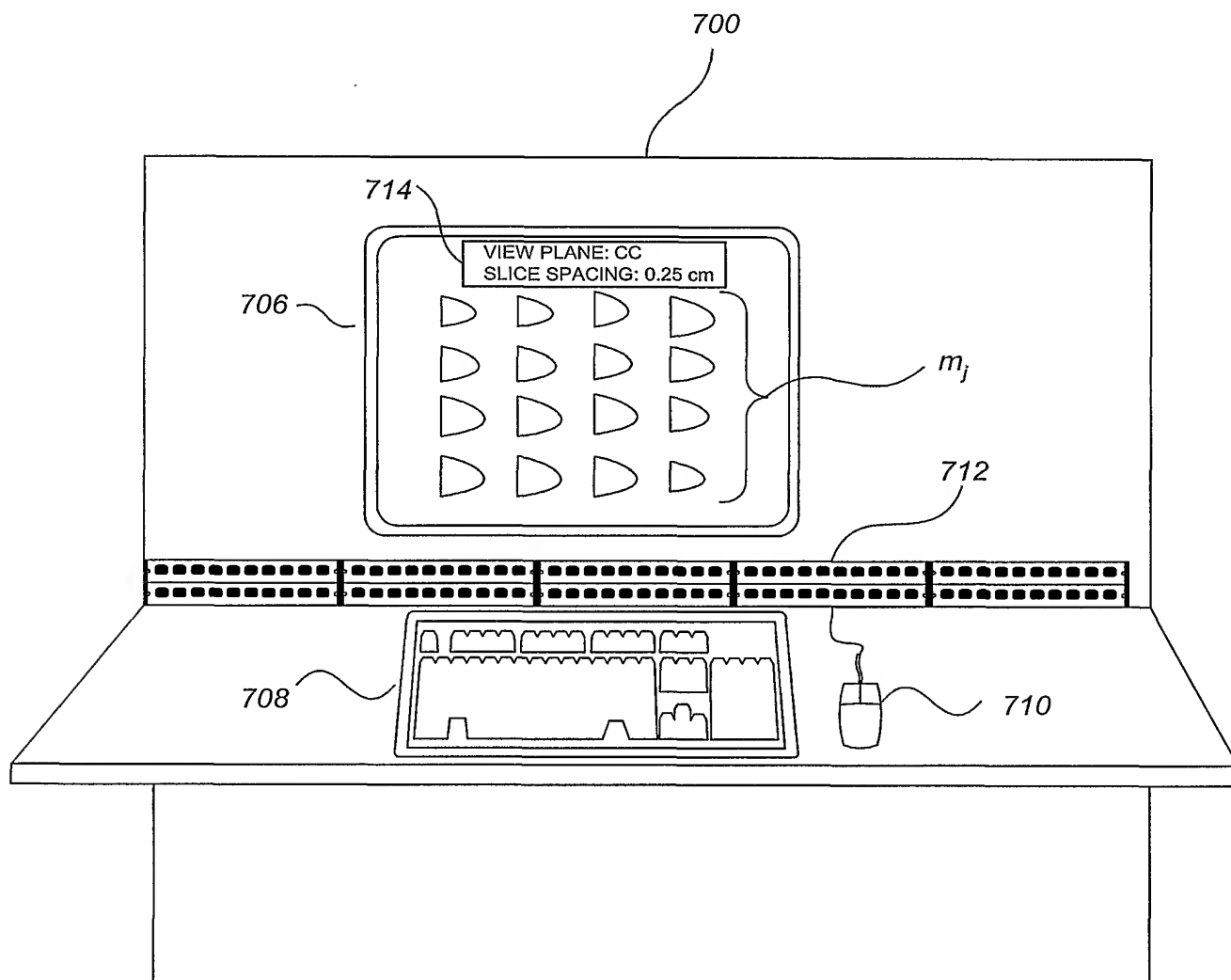


FIG. 5

**FIG. 6**

**FIG. 7**



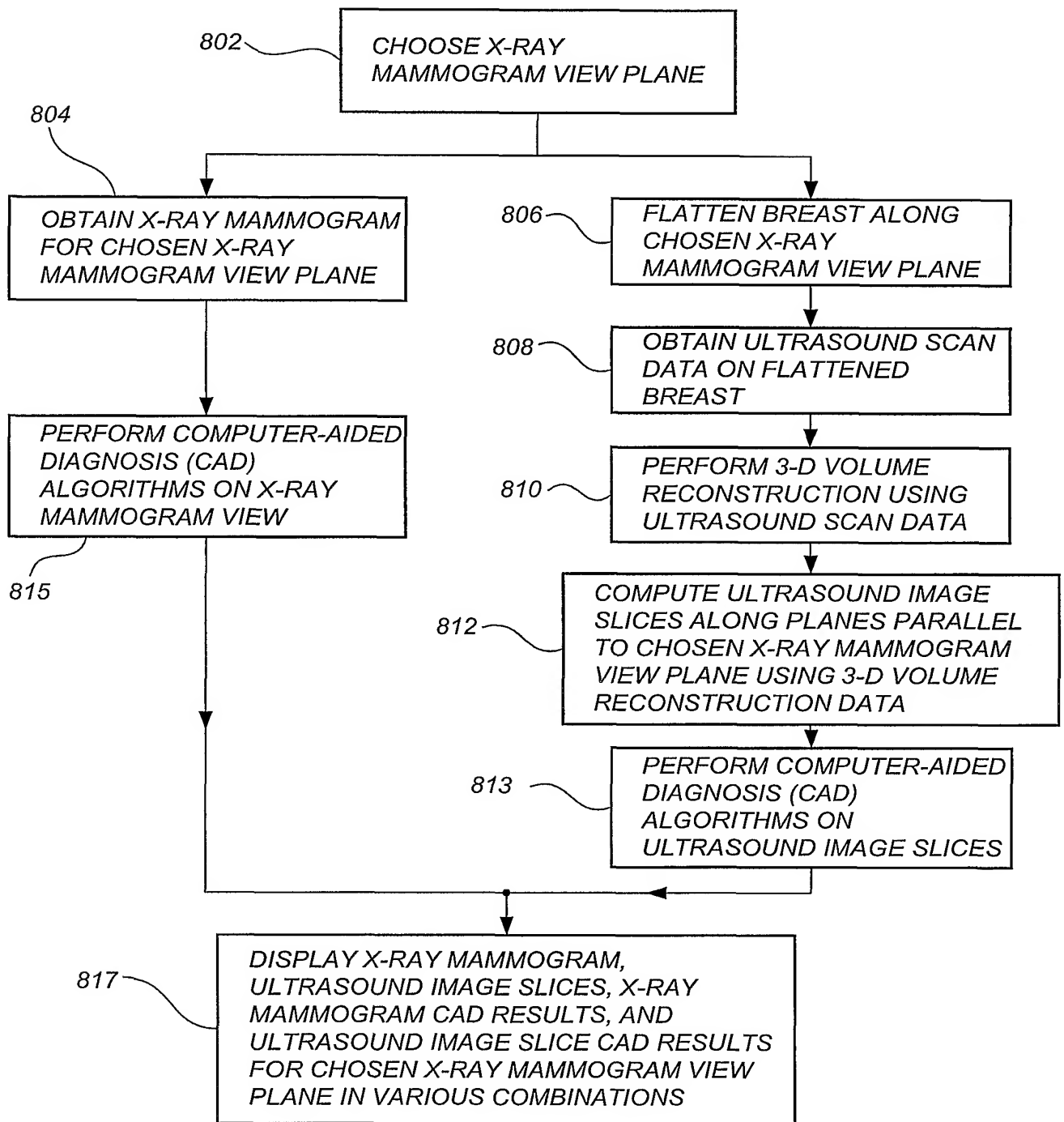


FIG. 8

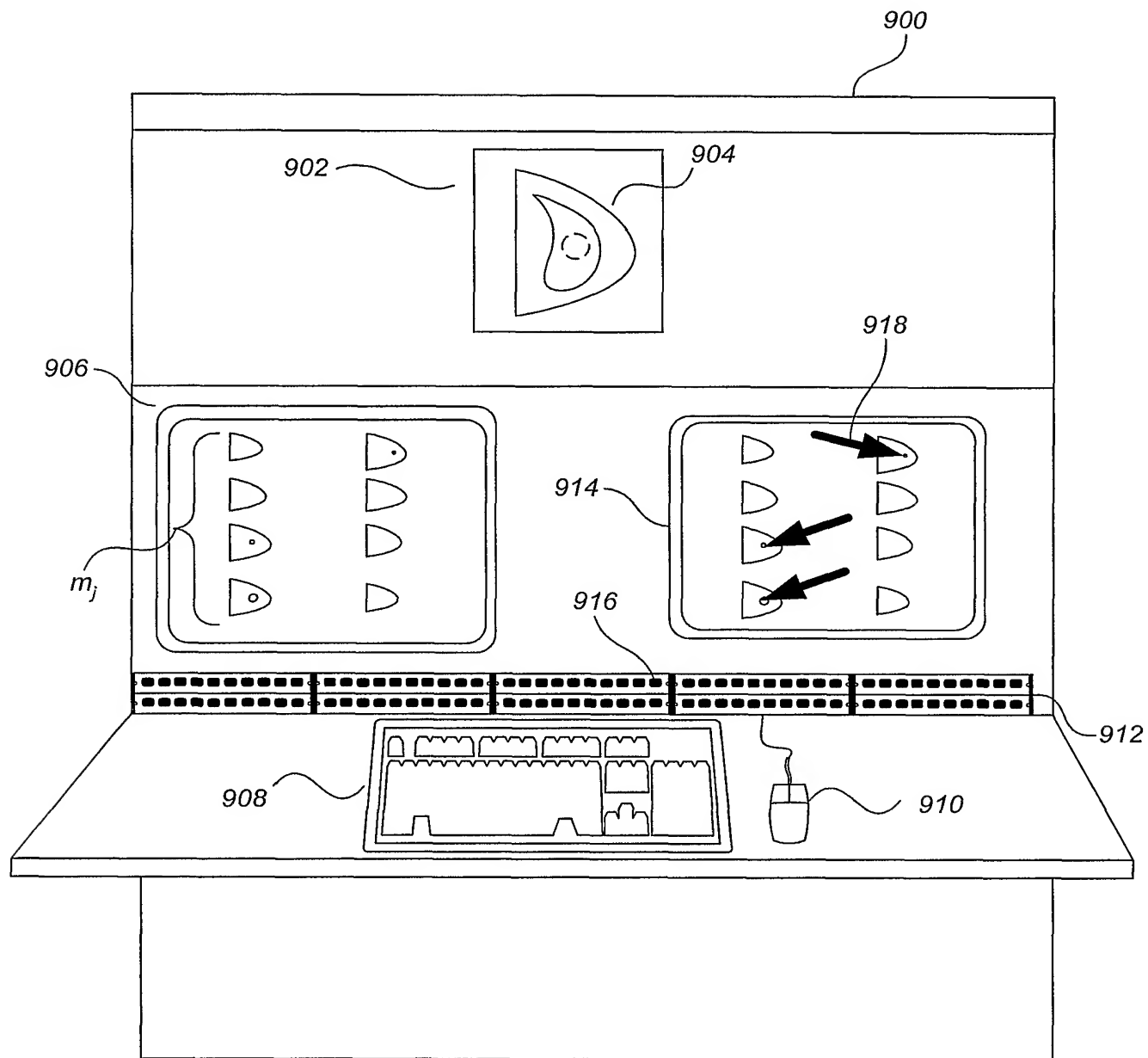


FIG. 9

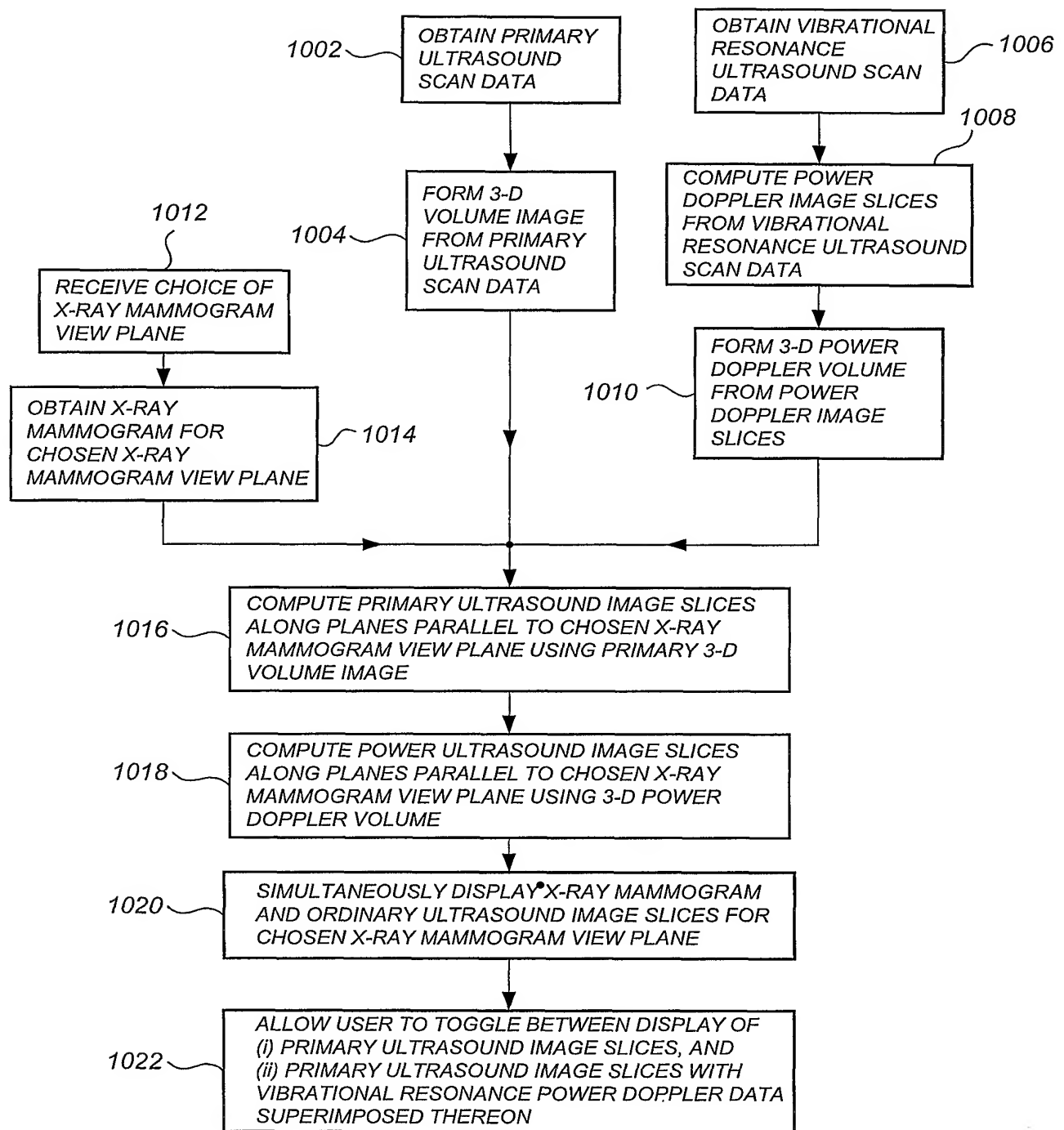


FIG. 10

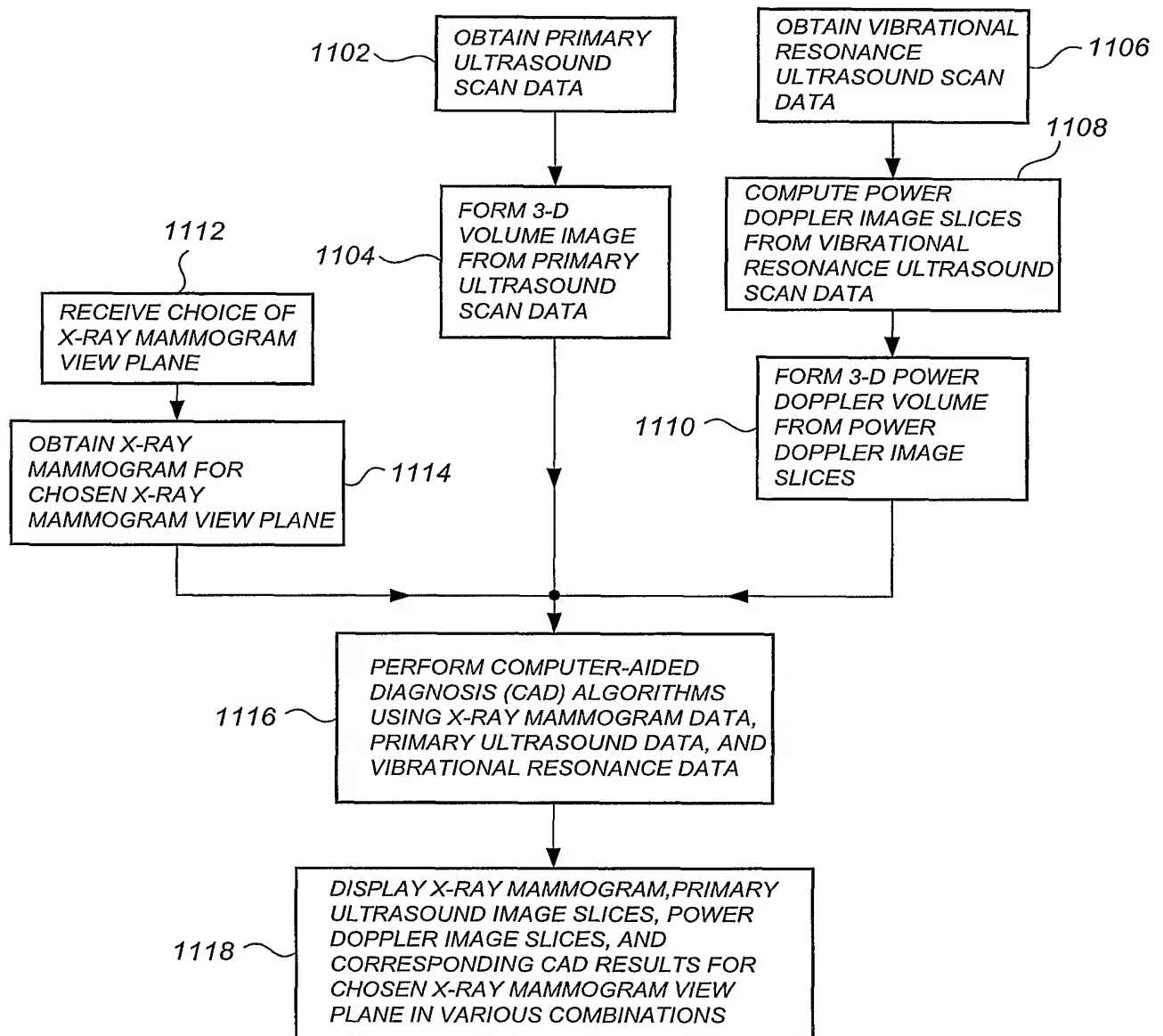


FIG. 11

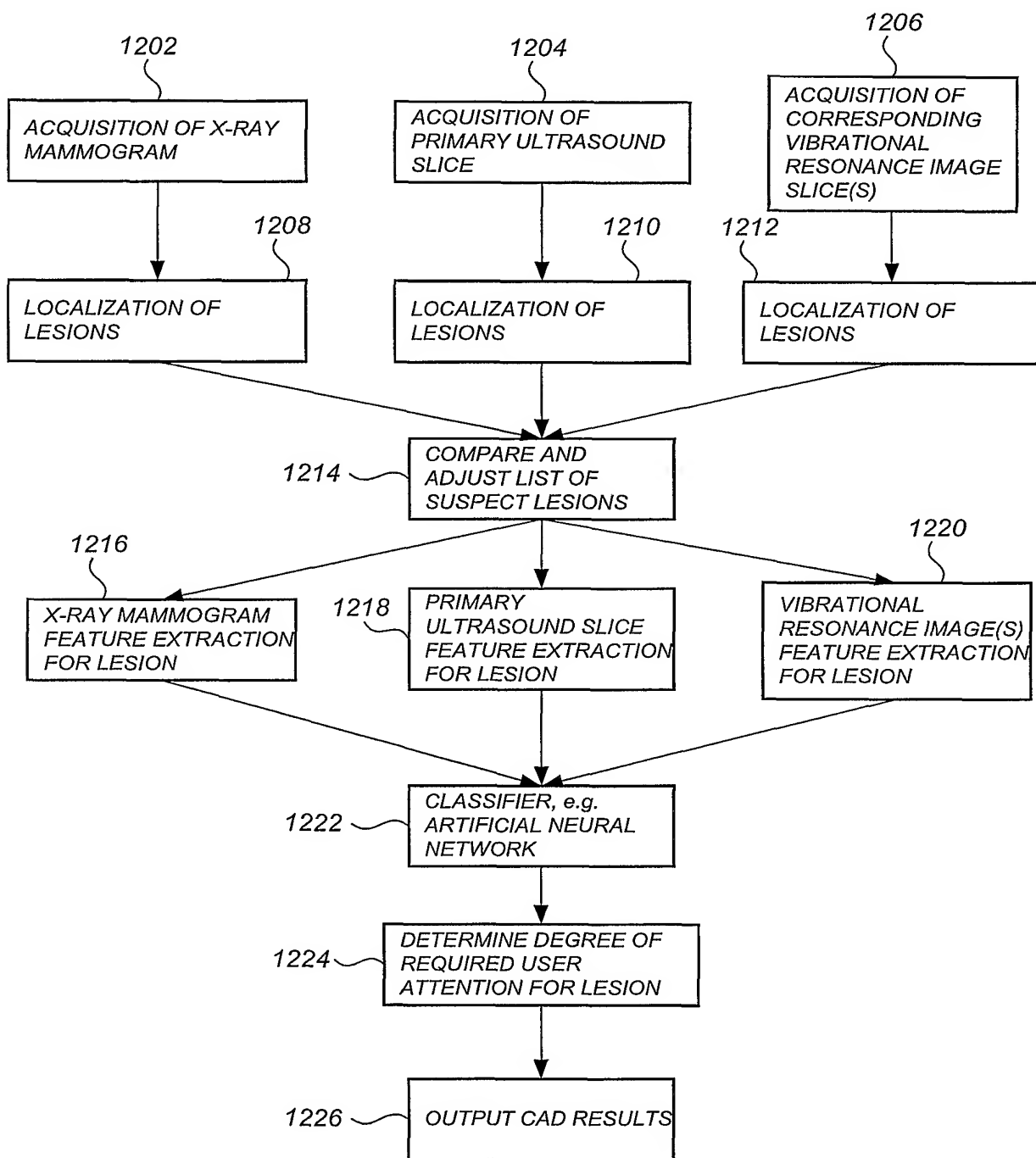


FIG. 12A

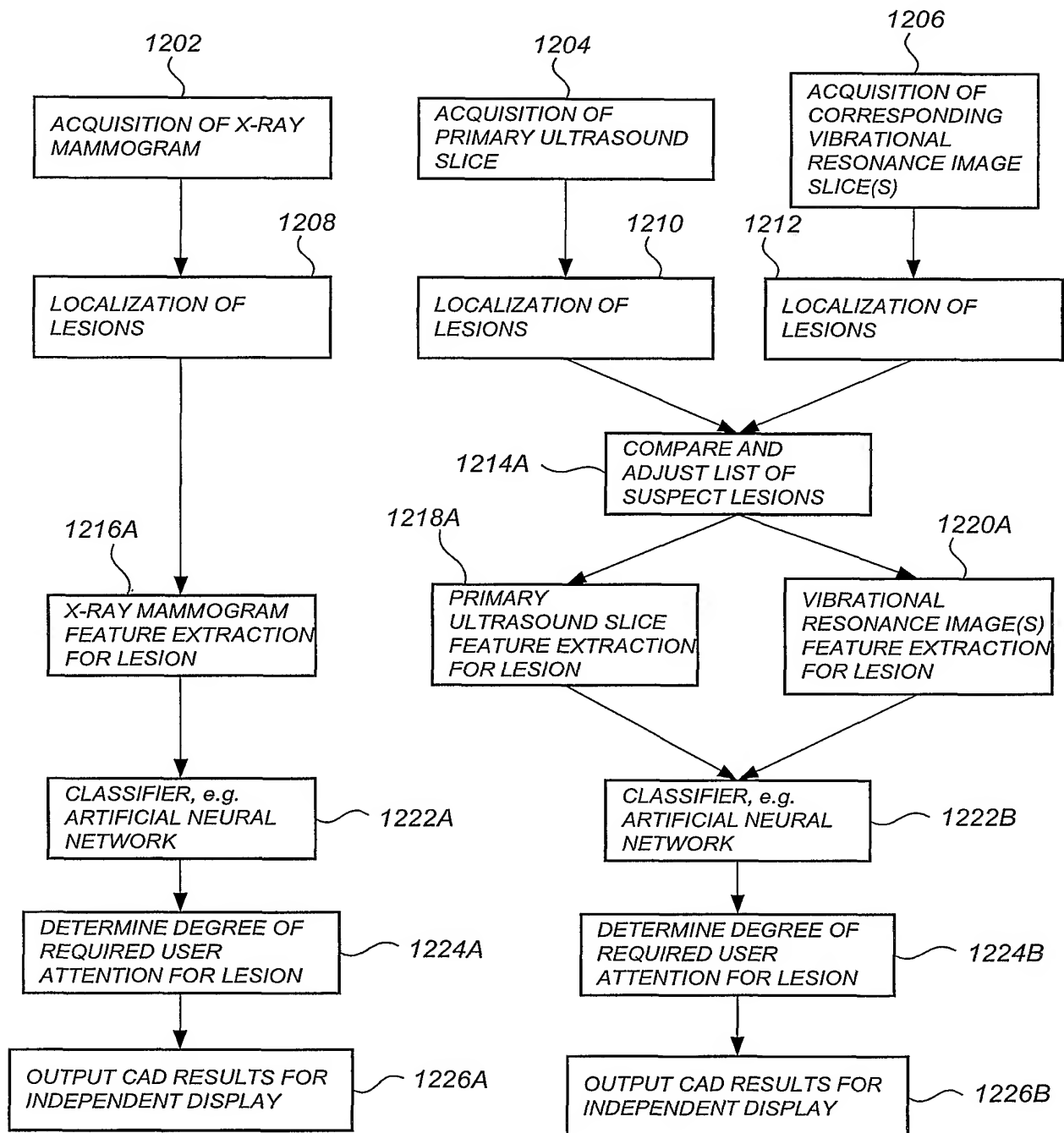


FIG. 12B

FIG. 13

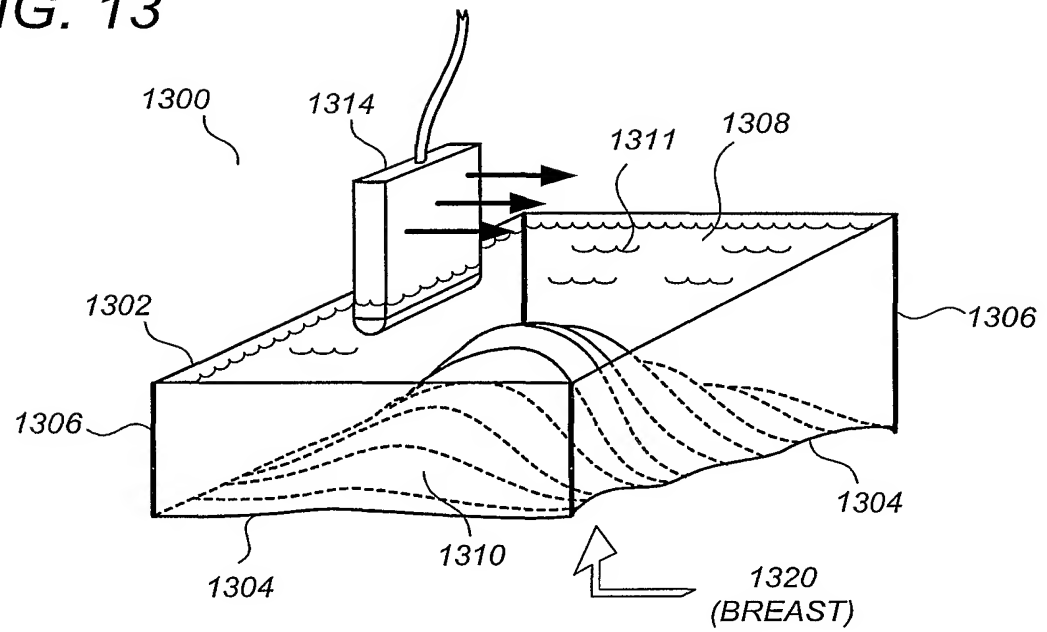


FIG. 14

